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Pred. No. is the number of results predicted by chance to have a

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and is derived by analysis of the total score distribution.

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6	17	94.4	18	6	AX104114 Sequence
7	17	94.4	18	6	AX353358 Sequence
8	17	94.4	24	6	AX463126 Sequence
9	13.2	73.3	37	6	AX463127 Sequence
10	13.2	73.3	39	6	AR089858 Sequence
11	13.2	73.3	37	6	AR167631 Sequence
12	12.8	71.1	48	6	AR151177 Sequence
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35	11.8	65.6	25	6	AX027188 Sequence
36	11.8	65.6	26	6	AX027200 Sequence
37	11.8	65.6	35	6	AR105706 Sequence
38	11.8	65.6	35	6	AX155967 Sequence
39	11.8	65.6	36	6	AX361817 Sequence
40	11.8	65.6	39	6	I26200 Sequence
41	11.8	65.6	40	6	AR182122 Sequence 34
42	11.8	65.6	43	3	AR169552 Sequence
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ALIGNMENTS

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ACCESSION
VERSION
KEYWORDS
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ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL

A89789
Sequence 11 from Patent WO9832462.
A89789
A89789.1 GI:6738303
unidentified.
unclassified.
1 (bases 1 to 18)
Lipford,G.B. and Heeg,K.
PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
Patent: WO 9832462-A 11 30-JUL-1998;

18 bp
DNA
linear
PAT 22-JAN-2000

Thu Dec 12 07:53:25 2002

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    1 (bases 1 to 18)
    Heeg,K.P. and Lipford,G.B.
    Pharmaceutical composition comprising a polynucleotide and an
    antigen especially for vaccination
    Patent: EP 0855184-A 11 29-JUL-1998;
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    Hartmann,G.D., Bratzler,R.L. and Krieg,A.U.
    Methods related to immunostimulatory nucleic acid-induced
    interferon
    Patent: WO 0122990-A 46 05-APR-2001;
    JOURNAL
    Coley Pharmaceutical Group, Inc. (US) ; UNIVERSITY OF IOWA RESEARCH
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    Bauer,S., Lipford,G. and Wagner,H.
    Process for high throughput screening of cpg-based
    immuno-agonist/antagonist
    Patent: WO 0222809-A 32 21-MAR-2002;
    JOURNAL
    Coley Pharmaceutical GmbH (DE)
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  REFERENCE
    1 (bases 1 to 18)
    Krieg,A.M., Schetter,C. and Vollmer,J.C.
    Immunostimulatory nucleic acids
    Patent: WO 0122972-A 306 03-APR-2001;
    JOURNAL
    UNIVERSITY OF IOWA RESEARCH FOUNDATION (US) ; Coley Pharmaceutical
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KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE
AUTHORS Weiner,G. and Hartmann,G.
TITLE Methods for enhancing antibody-induced cell lysis and treating
JOURNAL Patent: WO 0197843-A 386 27-DEC-2001.
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DEFINITION Sequence 9 from Patent WO0250108.
ACCESSION AX463126
VERSION AX463126.1 GI:21886107
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE
AUTHORS Marchal,G., Pescher,P. and Romain,F.
TITLE Immunogenic glycopeptides, screening, preparation and uses
JOURNAL Patent: WO 0250108-A 9 27-JUN-2002;
PASTEUR INSTITUT (FR)
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ACCESSION AR167631
VERSION AR167631.1 GI:17903423
KEYWORDS
SOURCE unknown.
ORGANISM unknown.
REFERENCE
AUTHORS Lee,F., Huszar,D. and Gu,W.
TITLE Screening methods for compounds useful in the regulation of body weight
JOURNAL Patent: US 6287763-A 41 11-SEP-2001;
FEATURES
Location/Qualifiers

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VERSION AX463127.1 GI:21886108
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ORGANISM
REFERENCE
AUTHORS Marchal,G., Pescher,P. and Romain,F.
TITLE Immunogenic glycopeptides, screening, preparation and uses
JOURNAL Patent: WO 0250108-A 10 27-JUN-2002;
PASTEUR INSTITUT (FR)
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DEFINITION Sequence 140 from patent US 5994075.
ACCESSION AR089858
VERSION AR089858.1 GI:10016613
KEYWORDS
SOURCE unknown.
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REFERENCE
AUTHORS Goodfellow,P.N.
TITLE Methods for identifying a mutation in a gene of interest without a
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JOURNAL Patent: US 5994075-A 140 30-NOV-1999;
PASTEUR INSTITUT (FR)
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DEFINITION Sequence 41 from patent US 6287763.
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VERSION AR167631.1 GI:17903423
KEYWORDS
SOURCE unknown.
ORGANISM unknown.
REFERENCE
AUTHORS Lee,F., Huszar,D. and Gu,W.
TITLE Screening methods for compounds useful in the regulation of body weight
JOURNAL Patent: US 6287763-A 41 11-SEP-2001;
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ACCESSION AR151177
VERSION AR151177.1 GI:15117227
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 48)
AUTHORS Marchionni, M. Andrew. and Johnson, C. D.
TITLE Homology cloning.
JOURNAL Patent: US 6232061-A 3 15-MAY-2001;
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ACCESSION AX103935
VERSION AX103935.1 GI:13920132
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 19)
AUTHORS Krieg, A. M., Schetter, C. and Vollmer, J. C.
TITLE Immunostimulatory nucleic acids
JOURNAL Patent: W0 0122972-A 127 05-APR-2001;
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LOCUS
DEFINITION Sequence 62 from patent US 6291173.
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ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS 1 (bases 1 to 19)
TITLE Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
JOURNAL Robbins, J. M. and Ritz, R.
FEATURES Ribozyne therapy for the treatment of proliferative skin and eye
diseases
Patent: WO 0130362-A 2667 03-MAY-2001;
IMMUSOL, INC. (US)
LOCATION/Qualifiers
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DEFINITION Sequence 413 from Patent W00197843.
ACCESSION AX355385
VERSION AX355385.1 GI:18620053
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1
AUTHORS Weiner, G. and Hartmann, G.
TITLE Methods for enhancing antibody-induced cell lysis and treating
cancer
JOURNAL Patent: WO 0197843-A 413 27-DEC-2001;
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US)
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REFERENCE   1 (bases 1 to 41)
AUTHORS     Bartel,P.L. and Tavtigian,S.V.
TITLE       WMS2--an MMAC1 interacting protein
JOURNAL     Patent: US 6291173-A 62 18-SEP-2001;
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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7	20	100.0	21	20 AAV82453	API comp oligonuc
8	20	100.0	21	21 AAZ8651	Rabbit AP-1 bindin
9	20	100.0	21	22 AAH26603	AP-1 oligonucleoti

10	20	100.0	21	22	AAH26603	Rat AP-1 synthetic
11	20	100.0	21	22	AAF87560	Consensus binding
12	20	100.0	21	24	ABL56586	Oligonucleotide sp
13	18.4	92.0	21	18	AAH85833	API oligonucleotide
14	18.4	92.0	21	20	AAZ25683	Transcription fact
15	18.4	92.0	21	20	AAH60221	Oligonucleotide AP
16	18.4	92.0	21	20	AAH76049	cAMP response elem
17	18.4	92.0	21	20	AAV08337	CRE element coding
18	18.4	92.0	21	21	AAA52331	AP-1 footprinting
19	18.4	92.0	21	22	AAI70580	Transcription fact
20	18.4	92.0	21	22	AAH13506	Rat AP-1 synthetic
21	18.4	92.0	21	22	AAF85085	Probe AP-1 used in
22	18.4	92.0	21	24	ABK97978	Cell-TRAP method a
23	18.4	92.0	21	24	ABK97979	Cell-TRAP method a
24	18.4	92.0	21	24	ABK98231	Nucleic acid bindi
25	18.4	92.0	21	24	ABK98232	Nucleic acid bindi
26	18.4	92.0	21	24	ABL60768	Nuclear factor-kap
27	16.8	84.0	21	19	AAV44855	Probe for AP-1 gen
28	16.8	84.0	21	22	AAH26605	AP-1 mutant oligon
29	16.8	84.0	21	22	AAH13505	Rat AP-1 synthetic
30	15.2	76.0	21	22	AAH13507	Rat AP-1 synthetic
31	15.2	76.0	21	24	ABL60767	Nuclear factor-kap
32	14.4	72.0	19	24	ABL44268	Human chromosome 1
33	14.4	72.0	22	15	AAQ67303	Detection probe fo
34	14.4	72.0	22	21	AAZ56897	AP-1 consensus seq
35	14.4	72.0	25	22	AAF99633	Immunostimulatory
36	14.4	72.0	25	22	AAF99634	Immunostimulatory
37	14.4	72.0	25	24	ABL38910	Immunostimulatory
38	14.4	72.0	25	24	ABL38913	Immunostimulatory
39	14.4	72.0	60	24	ABN34717	Human spliced tran
40	14.2	71.0	60	24	ABN36654	Human spliced tran
41	13.8	69.0	21	16	AAQ97424	Phage lambda (304-
42	13.8	69.0	21	22	AAI70282	Phage lambda genom
43	13.8	69.0	25	24	AAH42574	Phospholipase A1 p
44	13.8	69.0	47	21	AAZ68783	Human map-related
45	13.8	69.0	60	22	AAF82009	1.0 kb DNA fragmen

ALIGNMENTS

RESULT 1
AAV46003
ID AAV46003 standard; DNA; 20 BP.
XX
AC AAV46003;
XX
DT 16-OCT-1998 (first entry)
XX
DE Immune adjuvant AP-1 #2.
XX
KW Immune system; adjuvant; vaccine; cancer; prophylactic; pathogenicity;
KW modulator; tolerance; regulator; helper cell; antigen; immunoglobulin;
KW Ig class; autoimmune response; T-cell; B-cell; tumour; ss.
XX
OS Class Bacteria.
XX
PN EP855184-A1.
XX
PD 29-JUL-1998.
XX
PF 23-JAN-1997; 97EP-0101019.
XX
PR 23-JAN-1997; 97EP-0101019.
XX
PA (HEG/) HEG K.
PA (LIPF/) LIPFORD G B.
PA (WAGN/) WAGNER H.
PI Heeg K, Lipford GB, Wagner H;
XX
WPI; 1998-389630/34.
XX

PT Antigenic composition comprises polynucleotide fragment and antigen
PT - used as vaccine to treat or prevent e.g. cancer or pathogen
PT infections and to modulate immune response e.g. tolerance break and
PT regulation of TH1/TH2 cells
XX
PS Example 5; Page 8; 28pp; English.
XX
CC AAV45993-V46019 are fragments of bacterial polynucleotides which are
CC used as immune adjuvants for inclusion into vaccines to treat cancer and
CC for prophylaxis and/or treatment of conditions caused by pathogenic
CC micro-organisms. The polynucleotide is used for modulation of an immune
CC response and the modulation is selected from the group break of
CC tolerance, regulation of TH1/TH2 helper cell responses, switch of Ig
CC classes, treatment of autoimmune responses and induction of tolerances.
CC DNA oligomers are used to enhance the reactivity of immune cells to
CC viral, bacterial and parasitic antigens, to break tolerance in anergic T
CC and B cells e.g. against tumour antigens, as adjuvants in vaccination
CC against tumour-defined antigens and immunostimulatory substances in an
CC immune response against tumours and to suppress immune reactions of the
CC innate and acquired immune system. The composition is inexpensive and
CC stable and does not cause lethal shock, which happens with prior art
CC bacterial sequences.
XX
SQ Sequence 20 BP; 5 A; 5 C; 6 G; 4 T; 0 other;
Query Match 100.0%; Score 20; DB 19; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.43;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GCTTGATGACTACGCCGAA 20
DB 1 GCTTGATGACTACGCCGAA 20
RESULT 2
AAZ48028
ID AAZ48028 standard; DNA; 20 BP.
XX
AC AAZ48028;
XX
DT 08-MAR-2000 (first entry)
XX
DE Immune remodeling inducing CpG oligonucleotide control SEQ ID NO:108.
XX
KW Haematopoiesis; regulation; CpG oligonucleotide; phosphorothioate;
KW immune remodeling; thrombopoiesis; anaemia; immune system; cancer;
KW immune response; allergic reaction; infectious disease; asthma;
KW thrombocytopaenia; immunohaemolytic disorder; genetic disorder;
KW haemoglobinopathy; kidney failure; chronic inflammatory disorder;
KW rheumatoid arthritis; ss.
XX
OS Synthetic.
XX
PN WO958118-A2.
XX
PD 18-NOV-1999.
XX
PF 14-MAY-1999; 99WO-IB01285.
XX
PR 14-MAY-1998; 98US-0085516.
XX
PR 02-FEB-1999; 99US-0241653.
XX
PA (CPGI-) CPG IMMUNOPHARMACEUTICALS GMBH.
XX
PA (CPGI-) CPG IMMUNOPHARMACEUTICALS INC.
XX
PI Wagner H, Lipford G;
XX
DR WPI; 2000-062261/05.
XX
PT Use of CpG containing oligonucleotides for, e.g. inducing an
PT antigen-specific immune response
XX
PS Example 1; Page 51; 116pp; English.

XX The present invention describes a method using CpG containing
CC oligonucleotides (ONs) for regulating immune system remodeling and for
CC regulating haematopoiesis. The method for inducing an antigen-specific
CC immune response comprises: (1) administering an ON having a sequence
CC including at least the formula (1); and (2) exposing the subject to an
CC antigen at least 3 days after the ON is administered to the subject to
CC produce an antigen-specific immune response: 5' XICGX2 3' (1), where
CC the ON = includes at least 8 nucleotides; C and G = unmethylated, and
CC X1 and X2 = nucleotides. The method can be used for inducing an immune
CC response against an antigen such as cells, cell extracts, proteins,
CC polysaccharides, polysaccharide conjugates, lipids, glycolipids,
CC carbohydrates, viral extracts, viruses, bacteria, fungi, parasites and
CC allergens. It can be used in a subject at risk of developing cancer or
CC an allergic reaction. It can also be used for treating an infectious
CC disease, allergic diseases and asthma, as well as thrombocytopaenia
CC which is drug-induced, due to an autoimmune disorder such as idiopathic
CC thrombocytopenic purpura, or resulting from accidental or therapeutic
CC radiation exposure. It can also be used for treating anaemia such as
CC drug-induced anaemia, immunohaemolytic disorder, genetic disorders such
CC as haemoglobinopathy and inherited haemolytic anaemia, inadequate
CC production of adequate iron stores, chronic disease such as kidney
CC failure, and chronic inflammatory disorder such as rheumatoid arthritis,
CC or anaemia resulting from accidental or therapeutic radiation exposure.
CC AAZ47932 to AAZ48029 represent phosphorothioate CpG oligonucleotides
XX used in the exemplification of the present invention.
SQ Sequence 20 BP; 5 A; 5 C; 6 G; 4 T; 0 other;
Query Match 100.0%; Score 20; DB 21; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.43;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GCTTGATGACTACGCCGAA 20
DB 1 GCTTGATGACTACGCCGAA 20
RESULT 3
AAL39184
ID AAL39184 standard; DNA; 20 BP.
XX
AC AAL39184;
XX
DT 05-SEP-2002 (first entry)
XX
DE Murine Toll-like receptor related CpG DNA SEQ ID No 59.
XX
KW Murine Toll-like receptor; TLR9; TLR7; TLR8; ISNA; ds.
XX
OS Unidentified.
XX
PN WO200222809-A2.
XX
PD 21-MAR-2002.
XX
PF 17-SEP-2001; 2001WO-US29229.
XX
PR 15-SEP-2000; 2000US-233035P.
XX
PR 23-JAN-2001; 2001US-263657P.
XX
PR 17-MAY-2001; 2001US-291726P.
XX
PR 22-JUN-2001; 2001US-300210P.
XX
PA (COLE-) COLEY PHARM GMBH.
XX
PA Bauer S, Lipford G, Wagner H;
XX
DR WPI; 2002-393964/42.
XX
PT New isolated murine Toll-like receptor (TLR)9, TLR7, TLR8 polypeptides,
PT useful for identifying species specificity of immunostimulatory nucleic
PT acid and identifying immunostimulatory nucleic acids
XX

PS Disclosure: Page 76; 195pp; English.

XX The invention relates to isolated murine Toll-like receptors (TLR)9.

CC TLR7 and TLR8 polypeptides. These polypeptides comprise fully defined

CC sequences of 1032, 1050 or 1032 amino acids as given in specification, or

CC their fragments, where TLR9, TLR7 and TLR8 polypeptides or their

CC fragments have an amino acid sequence which is identical to human TLR9,

CC TLR7 or TLR8 polypeptides or their fragment except for at least one amino

CC acid of a murine TLR polypeptide. The isolated nucleic acids of the

CC invention are useful for inhibiting TLR9 signalling activity in a cell.

CC TLR7, TLR8 and TLR9 polypeptides are useful for identifying nucleic acid

CC molecules which interact with a TLR polypeptide or its fragment. The

CC TLR7, TLR8 or TLR9 polypeptides are also useful for identifying ISNA. The

CC TLR7, TLR8 and TLR9 polypeptides are also useful for comparing TLR9

CC signalling activity of a test compound (that is not a nucleic acid, and

CC is a polypeptide or a part of a combinatorial library of compounds) with

CC an ISNA. The TLR7, TLR8 and TLR9 polypeptides are also useful for

CC identifying species specificity of an ISNA. The isolated nucleic acids of

CC the invention are useful as probes or primers. This polynucleotide

CC sequence represents DNA relating to the isolated Toll-like receptors of

CC the invention.

XX

SQ Sequence 20 BP; 5 A; 5 C; 6 G; 4 T; 0 other;

Query Match 100.0%; Score 20; DB 24; Length 20;

Best Local Similarity 100.0%; Pred. No. 0.43;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20

DB 1 GCTTGATGACTCAGCGGAA 20

|||||

RESULT 4

AAV44856

ID AAV44856 standard; DNA; 21 BP.

XX

AC AAV44856;

XX

DT 21-OCT-1998 (first entry)

XX

DE Probe for AP-1 gene.

XX

KW Entry mediator gene; herpesvirus; HVEM; tumour necrosis factor receptor;

KW gene expression regulator; cellular stress; inflammatory response;

KW lymphocyte activity regulator; autoimmune response; probe; ss.

XX

OS Synthetic.

OS Homo sapiens.

XX

PN WO9825967-A1.

XX

PD 18-JUN-1998.

XX

PF 05-DEC-1997; 97WO-US22278.

XX

PR 12-DEC-1996; 96US-0032705.

XX

PA (GETH) GENENTECH INC.

XX

PI Ashkenazi AJ, Marsters SA;

XX

DR WPI; 1998-348457/30.

XX

PT Herpesvirus entry mediator polypeptide, HVEM - useful, e.g. in

PT assays for HVEM and to produce antibodies and transgenic animals,

PT e.g. for drug screening

XX

PS Example 3; Page 32; 46pp; English.

XX

CC This sequence is a probe for AP-1, used to determine if the

CC herpesvirus entry mediator (HVEM) protein of the invention activates

CC NF-kappaB. The HVEM protein is useful in quantitative diagnostic assays

CC

CC for HVEM, in affinity purification of HVEM from recombinant cells/natural

CC sources and in competitive-type receptor binding assays. It can be used

CC to generate antibodies, also useful in diagnostic assays for HVEM and

CC affinity purification of HVEM. HVEM is believed to be a member of the

CC tumour necrosis factor receptor family. Transient transfection of HVEM

CC into human 293 cells caused marked activation of certain transcription

CC factors, suggesting that HVEM is involved in regulating gene expression

CC in response to infectious stimuli and cellular stress. The predominant

CC expression of HVEM mRNA in lymphocyte-rich tissues (e.g. spleen and

CC peripheral blood) suggests it may be a receptor in regulating lymphocyte

CC activity. Antibodies produced may be useful therapeutically, e.g.

CC antagonistic antibodies may be used to block excessive

CC inflammatory/autoimmune response resulting from e.g. AP-1 induction,

CC whilst agonistic antibodies may enhance HVEM regulation of such

CC induction. The DNA may be used diagnostically, e.g. to determine if DNA

CC and/or RNA encoding HVEM is present in cells, and to prepare HVEM

CC polypeptide recombinantly. It is also useful to produce non-human

CC transgenic animals (e.g. mice or rats), especially knockout animals

CC containing cells with an altered gene encoding HVEM polypeptide. Such

CC animals are useful in the development and screening of therapeutically

CC useful reagents.

XX

SQ Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;

Query Match 100.0%; Score 20; DB 19; Length 21;

Best Local Similarity 100.0%; Pred. No. 0.43;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20

DB 2 GCTTGATGACTCAGCGGAA 21

|||||

RESULT 5

AAV39812

ID AAV39812 standard; DNA; 21 BP.

XX

AC AAV39812;

XX

DT 24-SEP-1998 (first entry)

XX

DE AP1 sequence A.

XX

KW Mouse type 10 collagen promoter; AP1 sequence A; mutation; MEF-2;

KW identification; morphogen; OP-1; osteogenic protein 1; human c-fos; ss.

XX

OS Synthetic.

OS Homo sapiens.

XX

PN WO9826069-A1.

XX

PD 18-JUN-1998.

XX

PF 11-DEC-1997; 97WO-US23211.

XX

PR 12-DEC-1996; 96US-0764522.

XX

PA (CREA-) CREATIVE BIOMOLECULES INC.

XX

PI Harada S, Rodan GA, Sampath KT;

XX

DR WPI; 1998-348526/30.

XX

PT Identifying morphogen analogues - by using DNA defining a

PT morphogen-responsive transcription activating element from a mouse

PT type X collagen promoter

XX

PS Claim 10; Page 61; 88pp; English.

XX

CC A method has been developed for identifying a compound that induces a

CC morphogen-mediated biological effect. The method comprises: (a) providing

CC a test cell comprising DNA defining a morphogen-responsive transcription

CC activating element (MRTAE), and, in operative association, a reporter

CC gene encoding a detectable gene product, the DNA, when present in a
 CC morphogen-responsive cell contacted with morphogen, serving to induce
 CC transcription of the reporter gene; (b) exposing the test cell to a
 CC candidate compound, and (c) detecting expression of the detectable gene
 CC product, the expression indicating the ability of the candidate compound
 CC to induce the morphogen-mediated biological effect. The present invention
 CC also describes: (1) a method for assessing whether a sample comprises a
 CC substance competent to bind to DNA; (2) a method for identifying a
 CC candidate compound that induces a morphogen-mediated biological effect;
 CC (3) a method for monitoring cell differentiation or tissue morphogenesis;
 CC and (4) a method for identifying a tissue responsive to a morphogen or
 CC analogue. The methods can be used for obtaining morphogen analogues. In
 CC particular they can be used for obtaining analogues of osteogenic protein
 CC 1 (OP-1) for the treatment of a metabolic bone disease, e.g. a disease
 CC characterised by osteopenia. Analogues can also be obtained for treating
 CC mammals afflicted with ischemic, ulcerative or inflammatory tissue
 CC damage, or with injury or deterioration of a morphogen-responsive tissue
 CC such as bone, liver, kidney, nerve, gastrointestinal tract lining, tooth
 CC dentin or periodontal tissue. The present sequence represents API
 CC sequence A from the present invention.

XX Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;

Query Match 100.0%; Score 20; DB 19; Length 21;
 Best Local Similarity 100.0%; Pred. No. 0.43;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20
 |||||
 Db 2 GCTTGATGACTCAGCGGAA 21

RESULT 6

AA83976
 ID AAX83976 standard; DNA; 21 BP.

XX AC AAX83976;

XX 08-SEP-1999 (first entry)

DE API sequence A.

XX Mouse; type 10 collagen promoter; API sequence A; osteogenic protein;
 KW OP-1; morphogen; bone morphogenic protein; BMP; soft tissue disorder;
 KW apoptosis; morphogen-activated regulatory pathway; tumour;
 KW cellular immune rejection; viral disease; ss.

XX Unidentified.

XX WO9931136-A2.

XX 24-JUN-1999.

PF 16-DEC-1998; 98WO-US26788.

XX 01-DEC-1998; 98US-0110498.

PR 17-DEC-1997; 97US-0069931.

XX (CREA-) CREATIVE BIOMOLECULES INC.

XX Cohen CM, Kawabata M, Miyazono K, Oeda E, Sampath KT;

XX WPI; 1999-418756/35.

XX Maintaining or restoring tissue-appropriate phenotype

XX Disclosure; Page 45; 50pp; English.

XX A method has been developed for maintaining or restoring tissue-
 CC appropriate phenotype by expression of a phenotype-specific protein or
 CC by inhibiting an intracellular pathway that induces expression of a
 CC gene that is an inhibitor of normal phenotype. The method is for
 CC restoring cellular phenotype in a cell effected by disease, damage or

CC age. The method comprises activating an intracellular pathway that
 CC induces expression of a phenotype-specific gene. Another method is also
 CC described for restoring cellular phenotype in a cell effected by
 CC disease, damage or age, comprising inhibiting an intracellular pathway
 CC that induces expression of a gene (especially TGF-beta) that is an
 CC inhibitor of normal phenotype. The methods can be used to treat soft
 CC tissue disorders by affecting apoptosis by modulating a morphogen-
 CC activated regulatory pathway e.g. in tumours, cellular immune rejection
 CC and viral diseases. The present sequence is used in the exemplification
 CC of the present invention.

XX Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;

Query Match 100.0%; Score 20; DB 20; Length 21;
 Best Local Similarity 100.0%; Pred. No. 0.43;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20
 |||||
 Db 2 GCTTGATGACTCAGCGGAA 21

RESULT 7

AAV82453
 ID AAV82453 standard; DNA; 21 BP.

XX AC AAV82453;

XX 12-APR-1999 (first entry)

DE API comp oligonucleotide used in competition analysis.

KW Vascular endothelial growth factor; VEGF; human; hypoxia;
 KW vascular disease; tumour; cancer; angiogenesis; wound healing;
 KW therapy; diagnosis; ds.

XX Synthetic.

OS Homo sapiens.

XX WO9856936-A1.

XX 17-DEC-1998.

PF 10-JUN-1998; 98WO-EP03517.

PR 10-JUN-1997; 97EP-0109418.

XX (PLAC) MAX PLANCK GES FOERDERUNG WISSENSCHAFTEN.

XX Damert A, Plate K, Risau W;

XX WPI; 1999-080911/07.

XX New recombinant DNA - contains sequence that regulates
 PT hypoxia-induced expression, used for, e.g. treatment and diagnosis
 PT of vascular disease

XX Example 6; Page 41; 80pp; English.

XX Oligonucleotides hVEGF, hVEGF 5' DEL, API1 and API2 (see
 CC AAV82449-52), and competitor oligonucleotides API comp, ATF comp
 CC and VI30 (see AAV82453-55) were used in electrophoretic mobility
 CC shift assays to determine which transcription factor(s) bind to
 CC the cis-acting element that is involved in the potentiation of
 CC hypoxia inducible factor 1 (HIF-1) mediated hypoxic induction
 CC of vascular endothelial growth factor (VEGF) gene regulatory
 CC sequences. Experiments were performed using normoxic or hypoxic
 CC C6 cell nuclear extracts. An API consensus binding site was shown
 CC to compete for DNA-protein-complex formation at potentiating
 CC sequences. The invention relates to recombinant DNA molecules
 CC comprising regulatory sequences of the VEGF gene, especially the
 CC 3' untranslated region (see AAV82439) and promoter (see AAV82440),
 CC being capable of modulating hypoxia inducible expression of a

CC heterologous DNA in vivo. Such recombinant DNA molecules, vectors,
 CC host cells and transgenic animals can be used to identify and
 CC develop compounds and methods for diagnosing, treating, preventing
 CC and/or delaying a vascular or tumour disease.
 XX
 SQ Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;

Query Match 100.0%; Score 20; DB 20; Length 21;
 Best Local Similarity 100.0%; Pred. No. 0.43;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCCGGAA 20
 ||||||||||||||||
 Db 2 GCTTGATGACTCAGCCGGAA 21

RESULT 8
 AA289651
 ID AA289651 standard; DNA; 21 BP.
 XX
 AC AA289651;
 XX
 DT 28-JUN-2000 (first entry)
 XX
 DE Rabbit AP-1 binding site, consensus DNA sequence.
 XX
 KW Transcription factor; AP-1; C/EBP; CCAAT/enhancer binding protein;
 KW cardiant; gene therapy; coronary angioplasty; stent implantation;
 KW ET-1 gene; smooth muscle cell; aorta carotis; treatment; rabbit;
 KW coronary heart disease; ds.
 XX
 OS Oryctolagus cuniculus.
 XX
 PN DE29916160-U1.
 XX
 PD 09-MAR-2000.
 XX
 PF 31-AUG-1999; 99WO-US20047.
 XX
 PR 25-MAY-1999; 99DE-1023892.
 XX
 PA (CARD-) CARDIOGENE GENTHERAPEUTISCHE SYSTEME AG.
 XX
 DR WPI; 2000-247751/22.
 XX
 CC Double stranded oligonucleotides targeting genes encoding AP-1, C/EBP
 PT and related transcription factors useful for treatment of coronary
 PT heart disease -
 XX
 PS Example 6; Page 37; 53pp; German.
 XX
 CC This invention describes novel double stranded nucleic acids, which bind
 CC specifically to transcription factors AP-1, C/EBP (CCAAT/Enhancer
 CC Binding Protein) or related transcription factors. The products of the
 CC invention have cardiant activity and are used in gene therapy. Patients
 CC who have had coronary angioplasty or stent implantation may be at risk
 CC from an increased expression of certain genes due to the physical weight
 CC of the devices used inducing a number of genes involved in the cell
 CC cycle. The AP-1 and C/EBP specific double stranded nucleic acids can be
 CC used to block activation of the ET-1 gene in smooth muscle cells of the
 CC aorta carotis. The nucleic acids are therefore useful in treatment of
 CC coronary heart diseases. This sequence represents a consensus AP-1
 CC binding site isolated from rabbit (Oryctolagus cuniculus) which is used
 CC to illustrate the method of the invention.
 XX
 SQ Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;

Query Match 100.0%; Score 20; DB 21; Length 21;
 Best Local Similarity 100.0%; Pred. No. 0.43;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCCGGAA 20
 ||||||||||||||||

Db 2 GCTTGATGACTCAGCCGGAA 21

RESULT 9
 AAH26603
 ID AAH26603 standard; DNA; 21 BP.
 XX
 AC AAH26603;
 XX
 DT 12-NOV-2001 (first entry)
 XX
 DE AP-1 oligonucleotide.
 XX
 KW Melanoma differentiation associated gene-7; Mda-7; human;
 KW promoter; neuroblastoma; astrocytoma; glioblastoma multiforme;
 KW cervical cancer; breast cancer; colon cancer; prostate cancer;
 KW osteosarcoma; chondrosarcoma; tumour; therapy; PCR primer;
 KW electrophoretic mobility shift assay; EMSA; AP-1;
 KW transcription factor; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200164921-A1.
 XX
 PD 07-SEP-2001.
 XX
 PF 28-FEB-2001; 2001WO-US06782.
 XX
 PR 29-FEB-2000; 2000US-0515369.
 XX
 PA (UYCO) UNIV COLUMBIA NEW YORK.
 XX
 PI Fisher PB, Madireddi MT;
 XX
 DR WPI; 2001-565508/53.
 XX
 CC Melanoma differentiation associated gene-7 promoter capable of
 PT treating cancer comprises directing transcription of heterologous
 PT coding sequence encoding tumour suppressor polypeptide positioned
 PT downstream, useful for treating cancer -
 XX
 PS Example 2; Page 70; 132pp; English.
 XX
 CC The present sequence is that of an AP-1 transcription factor
 CC oligonucleotide. This was used in electrophoretic mobility
 CC shift assays (EMSA) to examine transcription factor binding to
 CC the human melanoma differentiation associated gene-7 (mda-7)
 CC promoter (see AAH26595). Results demonstrated that cJUN/AP-1 and
 CC C/EBP-beta transcription factors bind to defined sites within the
 CC mda-7 promoter during the process of terminal differentiation in
 CC human melanoma cells. The invention provides recombinant
 CC expression constructs in which the mda-7 promoter is operably
 CC linked to a coding sequence encoding a tumour suppressor protein.
 CC A pharmaceutical composition including the recombinant expression
 CC construct is used in a claimed method of treating melanoma,
 CC neuroblastoma, astrocytoma, glioblastoma multiforme, cervical
 CC cancer, breast cancer, colon cancer, prostate cancer, osteosarcoma,
 CC chondrosarcoma or a cancer of the central nervous system.
 XX
 SQ Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;

Query Match 100.0%; Score 20; DB 22; Length 21;
 Best Local Similarity 100.0%; Pred. No. 0.43;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCCGGAA 20
 ||||||||||||||||
 Db 2 GCTTGATGACTCAGCCGGAA 21

RESULT 10
 AAD13504
 ID AAD13504 standard; DNA; 21 BP.

KW Nuclear transcription factor AP-1; R-flurbiprofen; S-flurbiprofen;
 KW R-arylpropionic acid; joint destruction; antirheumatic;
 XX rheumatic disease; ss.
 OS Synthetic.
 XX WO200224190-A1.
 PN
 XX
 PD 28-MAR-2002.
 XX
 XX 24-SEP-2001; 2001WO-EP11004.
 PF
 XX 25-SEP-2000; 2000DE-1047319.
 PR
 XX (PAZA-) PAZ ARZNEIMITTEL ENTWICKLUNGS GMBH.
 XX
 XX Geisslinger G, Groesch S;
 XX
 XX WPI; 2002-401951/43.
 DR
 XX Use of R-arylpropionic acids in the production of medicaments, useful
 PT for the treatment of illnesses, e.g. rheumatic diseases, influenced by
 PT activation of nuclear transcription factor -
 XX
 XX Example; Page 10; 20pp; German.
 XX
 XX The present sequence represents an oligonucleotide specific for nuclear
 CC transcription factor AP-1. The oligonucleotide was used to demonstrate
 CC that while R-flurbiprofen inhibited AP-1 DNA binding in a
 CC dosage-dependent manner (binding was completely suppressed at 1000
 CC microgram), S-flurbiprofen reduced AP-1 DNA binding at only 1000
 CC microgram). The specification describes the use of R-arylpropionic acids
 CC or their salts or derivatives in the production of medicaments which
 CC inhibit the activation of nuclear transcription factor AP-1 and are
 CC useful for the treatment of illnesses which are influenced by this
 CC factor. The R-arylpropionic acids arrest joint destruction but do not
 CC have the severe side effects associated with standard therapy using
 CC long-term antirheumatics. The medicaments are used in the treatment of
 CC rheumatic diseases, preferably in combination with basic therapeutics.
 XX
 XX Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;
 SQ
 Query Match 100.0%; Score 20; DB 24; Length 21;
 Best Local Similarity 100.0%; Pred. No. 0.43;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GCTTGATGACTCAGCCGGAA 20
 Db |||||||
 2 GCTTGATGACTCAGCCGGAA 21
 RESULT 13
 AAT85833
 ID AAT85833 standard; DNA; 21 BP.
 XX
 AC AAT85833;
 XX
 XX 21-NOV-1997 (first entry)
 DT
 XX
 DE AP1 oligonucleotide used in gel shift assay.
 XX
 KW Activating transcription factor 1; ATF1; CREB; recognition sequence;
 KW cyclic AMP responsive element binding protein; inhibition; binding;
 KW proliferation; virus; cancer; HTLV1; leukaemia; antibody; ss.
 XX
 OS Synthetic.
 XX
 XX US5641486-A.
 PN
 XX 24-JUN-1997.
 PD
 XX 18-MAR-1994; 94US-0210880.
 PF
 XX

PR 18-MAR-1994; 94US-0210880.
 XX
 XX (UYNE-) UNIV NEBRASKA.
 XX
 XX Hinrichs SH, Orten DJ;
 XX
 XX WPI; 1997-340900/31.
 DR
 XX Inhibiting replication of cancer cells or viruses - with inhibitor
 PT that binds to peptide sequence of activating transcription factor 1
 XX
 XX Example 2; Column 6; 17pp; English.
 XX
 CC This oligonucleotide sequence corresponds to the recognition sequence AP1
 CC to which members of the activating transcription factor 1 (ATF1)-cyclic
 CC AMP responsive element binding protein (CREB) family of proteins bind.
 CC The sequence was used in a gel shift mobility assay to identify agents
 CC which inhibit the binding of ATF1 to its recognition sequence. The
 CC agents are preferably antibodies, small molecules or polypeptides,
 CC especially the complementarity determining region of monoclonal antibody
 CC MAb4. The agents cause inhibition of transcription by dissociating ATF1
 CC from its target gene and thus will prevent proliferation of e.g. a virus
 CC or cancer cell, such as HTLV1-mediated leukaemic cell proliferation.
 XX
 XX Sequence 21 BP; 5 A; 5 C; 7 G; 4 T; 0 other;
 SQ
 Query Match 92.0%; Score 18.4; DB 18; Length 21;
 Best Local Similarity 95.0%; Pred. No. 3.1;
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 GCTTGATGACTCAGCCGGAA 20
 Db |||||||
 2 GCTTGATGACTCAGCCGGAA 21
 RESULT 14
 AAZ25683
 ID AAZ25683 standard; DNA; 21 BP.
 XX
 AC AAZ25683;
 XX
 XX 06-JAN-2000 (first entry)
 DT
 XX Transcription factor AP-1 (c-jun) oligonucleotide.
 DE
 XX Neuroprotective; calcium binding; stroke; neurodegenerative disease;
 KW blood-brain barrier; cerebral ischaemia; Alzheimer's disease;
 KW memory deficit; aging; ss.
 XX
 OS Synthetic.
 XX
 XX WO9940112-A1.
 PN
 XX 12-AUG-1999.
 PD
 XX 28-JAN-1999; 99WO-US01786.
 PF
 XX 10-FEB-1998; 98US-0021247.
 PR
 XX (NEUR-) NEUROMEDICA INC.
 PA
 XX Shashoua VE;
 PI
 XX WPI; 1999-610582/52.
 DR
 XX Neuroprotective peptides, which bind calcium, are useful for treating
 PT stroke and other neurodegenerative diseases -
 PT
 XX Example 4; Page 31; 65pp; English.
 PS
 XX The present invention describes a composition comprising an isolated
 CC peptide, which comprises the amino acid sequence (I) or (Ia):
 CC X1-XX3-XX5-XX7-XX9-XXX12 (I); X5-X6-X7-X8-X9-X10-X11-X12 (Ia); where

CC X1 = Asp, Gln, Gly or Tyr; X = any amino acid; X3 = Asp, Asn, Thr or Glu;
 CC X5 = Asp, Ser, Gly, Asn or Leu; X7 = Ala, Asp, Phe, Lys, Thr, Tyr, Arg,
 CC Val, Cys or Ser; X9 = Asp, Glu, Gly, Thr, Met or Asn; and X12 =
 CC Glu, Gln, Ala, Leu or Asn. (I) and (Ia) are neuroprotective calcium
 CC binding peptides. (I) is used to treat a condition characterized by
 CC cerebral ischaemia. (I) reduces the neurotoxic effect of cerebral
 CC ischaemia. (I) is used to increase neuronal cell AP-1 or NF-IL6
 CC transcription factor activity. The peptides are also useful for binding
 CC calcium. The peptide can be conjugated with a compound which facilitates
 CC transport across the blood brain barrier into the brain or it can be
 CC administered with a compound that increases transport across the blood
 CC brain barrier. Molecules that protect neurons against the ischaemic
 CC effects of stroke will also be useful for treating Alzheimer's disease,
 CC as well as the memory deficits that are characteristic of the aging
 CC process. The present sequence represents an oligonucleotide used in
 CC the exemplification of the present invention.
 XX
 SQ Sequence 21 BP; 5 A; 5 C; 7 G; 4 T; 0 other;
 Query Match 92.0%; Score 18.4; DB 20; Length 21;
 Best Local Similarity 95.0%; Pred. No. 3.1;
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 GCTTGATGACTCAGCGGAA 20
 ||||| ||||| ||||| |||||
 Db 2 GCTTGATGAGTCAGCGGAA 21
 RESULT 15
 AAX60221
 ID AAX60221 standard; DNA: 21 BP.
 XX
 AC AAX60221;
 XX
 DT 11-AUG-1999 (first entry)
 XX
 DE Oligonucleotide AP-1.
 KW keratin K1 based expression vector; epidermal cell expression;
 KW mammalian K1 Keratin gene; primer; ss.
 XX Synthetic.
 OS
 XX
 PN US5914265-A.
 XX
 PD 22-JUN-1999.
 XX
 PF 01-NOV-1993; 93US-0147777.
 XX
 PR 01-NOV-1993; 93US-0147777.
 PR 30-APR-1992; 92US-0876289.
 PR 29-OCT-1993; 93US-0145387.
 XX
 PA (BAYU) BAYLOR COLLEGE MEDICINE.
 PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX
 PI Greenhalgh DA, Roop DR, Rothnagel JA, Yuspa SH;
 XX
 DR WPI; 1999-370501/31.
 XX
 PT Keratin K1 expression vectors
 XX
 PS Example 16; Column 35; 50pp; English.
 XX
 CC The specification describes a keratin K1 based expression vector, which
 CC does not comprise a K1 keratin encoding gene sequence, for expression
 CC in an epidermal cell. The vector comprises a 5' flanking region from a
 CC mammalian gene including the necessary sequences for expression, a
 CC 3' flanking region from a mammalian K1 keratin gene which regulates
 CC expression in an epidermal cell, and a linker connecting the 5' flanking
 CC region to the nucleic acid sequence, where the linker has a position
 CC for inserting the nucleic acid sequence but lacks the coding sequence
 CC of a gene with which it is naturally associated. The genetic material

CC comprising the vector may encode a hormone, a receptor, a growth factor,
 CC an enzyme, a drug, a tumour suppressor, an apolipoprotein, a clotting
 CC factor, an antigen, an oncogene or a transforming gene. The present
 CC oligonucleotide was used in the course of the invention.
 XX
 SQ Sequence 21 BP; 5 A; 5 C; 7 G; 4 T; 0 other;
 Query Match 92.0%; Score 18.4; DB 20; Length 21;
 Best Local Similarity 95.0%; Pred. No. 3.1;
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
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 Db 2 GCTTGATGAGTCAGCGGAA 21
 Search completed: December 12, 2002, 01:36:26
 Job time : 105.319 secs

GenCore version 5.1.3
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OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:06:16 ; Search time 318.116 Seconds
(without alignments)
1829.698 Million cell updates/sec

Title: US-09-355-254F-12

Perfect score: 20

Sequence: 1 tcgatggggcgggcgagc 20

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0

Maximum DB seq length: 100

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

GenEmbl.*

- 1: gb.ba.*
- 2: gb.htg.*
- 3: gb.in.*
- 4: gb.om.*
- 5: gb.ov.*
- 6: gb.pat.*
- 7: gb.ph.*
- 8: gb.pl.*
- 9: gb.pr.*
- 10: gb.ro.*
- 11: gb.sts.*
- 12: gb.sy.*
- 13: gb.un.*
- 14: gb.vi.*
- 15: em.ba.*
- 16: em.fun.*
- 17: em.hum.*
- 18: em.in.*
- 19: em.mu.*
- 20: em.om.*
- 21: em.or.*
- 22: em.ov.*
- 23: em.pat.*
- 24: em.ph.*
- 25: em.pl.*
- 26: em.ro.*
- 27: em.sts.*
- 28: em.un.*
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- 30: em.htg_hum.*
- 31: em.htg_inv.*
- 32: em.htg_other.*
- 33: em.htg_mus.*
- 34: em.htg_pln.*
- 35: em.htg_rod.*
- 36: em.htg_rnam.*
- 37: em.htg_vrt.*
- 38: em_sy.*
- 39: em.htgo_hum.*
- 40: em.htgo_mus.*
- 41: em.htgo_other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match %	Length	DB ID	Description
1	20	100.0	20	6	A89791 Sequence 13
2	20	100.0	20	6	A90878 Sequence 13
3	20	100.0	20	6	AX023404 Sequence
4	20	100.0	20	6	AX455627 Sequence
5	20	100.0	22	6	AR060840 Sequence
6	20	100.0	22	6	AR070792 Sequence
7	20	100.0	22	6	AX195274 Sequence
8	20	100.0	48	6	AX377574 Sequence
9	20	100.0	48	6	AX377574 Sequence
10	20	100.0	49	6	AX127457 Sequence
11	20	100.0	49	6	AX127457 Sequence
12	20	100.0	49	6	AX127457 Sequence
13	20	100.0	49	6	AX127459 Sequence
14	20	100.0	49	6	AX377573 Sequence
15	19	95.0	21	6	AX104633 Sequence
16	19	95.0	21	6	AX104634 Sequence
17	19	95.0	21	6	AX355087 Sequence
18	19	95.0	21	6	AX355087 Sequence
19	19	95.0	21	6	AX355201 Sequence
20	19	95.0	46	6	I72381 Sequence 12
21	18.4	92.0	22	6	E07877 Synthetic n
22	18	90.0	46	6	I72381 Sequence 13
23	18	90.0	46	6	I72382 Sequence 12
24	16.8	84.0	49	6	AX377573 Sequence
25	15.8	79.0	31	6	AR091898 Sequence
26	15.8	79.0	31	6	E64477 Sugar-respo
27	15.8	79.0	31	6	I77207 Sequence 4
28	15.8	79.0	66	6	AX207285 Sequence
29	15.2	76.0	80	6	AX002555 Sequence
30	15.2	76.0	80	6	E27453 cdc25B Gene
31	14.8	74.0	27	6	AX182197 Sequence
32	14.8	74.0	27	6	AX382003 Sequence
33	14.4	72.0	58	6	AR034640 Sequence
34	14.4	72.0	58	6	I70120 Sequence 26
35	14.4	72.0	60	6	AR034639 Sequence
36	14.4	72.0	60	6	I70119 Sequence 25
37	13.6	68.0	33	6	AX363257 Sequence
38	13.6	68.0	59	9	HSGROUCH1
39	13.6	68.0	65	6	AR121168 Sequence
40	13.6	68.0	65	6	BD003583 Methods a
41	13.4	67.0	87	9	AF188738S5
42	13.2	66.0	39	6	AX006290 Homo sapi
43	13.2	66.0	51	6	AX204497 Sequence
44	13.2	66.0	81	14	AF221250 Hepatitis
45	13.2	66.0	81	14	AF221252 Hepatitis

ALIGNMENTS

RESULT 1	A89791	Sequence 13 from Patent WO9832462.	20 bp	DNA	linear	PAT 22-JAN-2000
LOCUS	A89791	Sequence 13 from Patent WO9832462.	20 bp	DNA	linear	PAT 22-JAN-2000
DEFINITION	A89791	Sequence 13 from Patent WO9832462.	20 bp	DNA	linear	PAT 22-JAN-2000
ACCESSION	A89791	Sequence 13 from Patent WO9832462.	20 bp	DNA	linear	PAT 22-JAN-2000
VERSION	A89791.1	GI:6738305				
KEYWORDS	unidentified.					
SOURCE	unidentified.					
ORGANISM	unclassified.					
REFERENCE	1 (bases 1 to 20)					
AUTHORS	Lipford, G.B. and Heeg, K.					
TITLE	PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND					
JOURNAL	OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION					
	Patent: WO 9832462-A 13 30-JUL-1998;					

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    LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
    Location/Qualifiers
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      /organism="unidentified"
      /db_xref="taxon:32644"
BASE COUNT
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Query Match
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Best Local Similarity
  100.0%; Pred. No. 3.3e+02;
Matches
  20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY
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DB
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  |||||

RESULT 2
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LOCUS
  A90878
DEFINITION
  Sequence 13 from Patent EP0855184.
ACCESSION
  A90878
VERSION
  A90878.1 GI:6739281
KEYWORDS
  unidentified.
SOURCE
  unidentified.
ORGANISM
  unclassified.
REFERENCE
  1 (bases 1 to 20)
AUTHORS
  Heeg,K.P. and Lipford,G.B.
TITLE
  Pharmaceutical composition comprising a polynucleotide and an
  antigen especially for vaccination
JOURNAL
  Patent: EP 0855184-A 13 29-JUL-1998;
  HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
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Matches
  20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY
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  |||||
DB
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  |||||

RESULT 3
AX023404
LOCUS
  AX023404
DEFINITION
  Sequence 19 from Patent WO0014217.
ACCESSION
  AX023404
VERSION
  AX023404.1 GI:10183804
KEYWORDS
  synthetic construct.
SOURCE
  synthetic construct
  artificial sequences.
ORGANISM
  1 (bases 1 to 20)
AUTHORS
  Lipford,G.B., Heeg,K. and Wagner,H.
TITLE
  G-motif oligonucleotides and uses thereof
JOURNAL
  Patent: WO 0014217-A 19 16-MAR-2000;
  LIPFORD GRAYSON B (DE); HEEG KLAUS (DE); WAGNER HERMANN (DE);
  CPG IMMUNOPHARMACEUTICALS GMBH (DE)
FEATURES
  source
    1..20
    /organism="synthetic construct"
    /db_xref="taxon:32630"
    /note="synthetic, no natural origin"
BASE COUNT
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Query Match
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Best Local Similarity
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Matches
  20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY
  1 TCGATCGGGCGGGCGGAGC 20
  |||||
DB
  1 TCGATCGGGCGGGCGGAGC 20
  |||||

RESULT 4
AX055627
LOCUS
  AX055627
DEFINITION
  Sequence 104 from Patent WO0222809.
ACCESSION
  AX055627
VERSION
  AX055627.1 GI:21714695
KEYWORDS
  synthetic construct.
SOURCE
  synthetic construct
  artificial sequences.
ORGANISM
  1
REFERENCE
  1
AUTHORS
  Bauer,S., Lipford,G. and Wagner,H.
TITLE
  Process for high throughput screening of cpq-based
  immuno-agonist/antagonist
JOURNAL
  Patent: WO 0222809-A 104 21-MAR-2002;
  Coley Pharmaceutical GmbH (DE)
FEATURES
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    /note="Synthetic oligonucleotide"
BASE COUNT
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Best Local Similarity
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Matches
  20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY
  1 TCGATCGGGCGGGCGGAGC 20
  |||||
DB
  1 TCGATCGGGCGGGCGGAGC 20
  |||||

RESULT 5
AR060640
LOCUS
  AR060640
DEFINITION
  Sequence 12 from patent US 5840832.
ACCESSION
  AR060640
VERSION
  AR060640.1 GI:5987090
KEYWORDS
  Unknown.
SOURCE
  Unknown.
ORGANISM
  1 (bases 1 to 22)
AUTHORS
  Ono,S,Jeremy. and Strominger,J.L.
TITLE
  Transcription factor regulating MHC expression, CDNA and genomic
  clones encoding same and retroviral expression constructs thereof
JOURNAL
  Patent: US 5840832-A 12 24-NOV-1998;
  Location/Qualifiers
FEATURES
  source
    1..22
    /organism="unknown"
BASE COUNT
  3 a 5 c 11 g 3 t
ORIGIN

Query Match
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Best Local Similarity
  100.0%; Pred. No. 3.2e+02;
Matches
  20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY
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  |||||
DB
  3 TCGATCGGGCGGGCGGAGC 22
  |||||

RESULT 6

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AR070792      AR070792      22 bp      DNA      linear      PAT 18-FEB-2000
LOCUS          Sequence 12 from patent US 5908762.
DEFINITION
ACCESSION      AR070792
VERSION        AR070792.1 GI:7221680
KEYWORDS
SOURCE         unknown.
ORGANISM       Unknown.
REFERENCE      1 (bases 1 to 22)
AUTHORS       Ono,S.Jeremy. and Strominger,J.L.
TITLE         transcription factor regulating MHC expression CDNA and genomic
              clones encoding same and retroviral expression constructs thereof
JOURNAL        Patent: US 5908762-A 12 01-JUN-1999;
FEATURES       Location/Qualifiers
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               /organism="unknown"
BASE COUNT    3 a      5 c      11 g      3 t
ORIGIN
Query Match   100.0%; Score 20; DB 6; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.2e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGATCGGGCGGGCGGAGC 20
   ||||||||||||||||||
Db 3 TCGATCGGGCGGGCGGAGC 22

RESULT 7
AX195274
LOCUS          AX195274      22 bp      DNA      linear      PAT 28-AUG-2001
DEFINITION     Sequence 10 from Patent WO0151671.
ACCESSION      AX195274
VERSION        AX195274.1 GI:15385825
KEYWORDS
SOURCE         synthetic construct.
ORGANISM       synthetic construct
              artificial sequences.
REFERENCE      1 (bases 1 to 22)
AUTHORS       McCarthy,J. and Cordell,B.
TITLE         Methods for identifying inhibitors of neuronal degeneration
JOURNAL        Patent: WO 0151671-A 10 19-JUL-2001;
              Scios Inc. (US)
FEATURES       Location/Qualifiers
               1..22
               /organism="synthetic construct"
               /db_xref="taxon:32630"
               /note="Synthetic Oligonucleotide"
BASE COUNT    3 a      5 c      11 g      3 t
ORIGIN
Query Match   100.0%; Score 20; DB 6; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.2e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGATCGGGCGGGCGGAGC 20
   ||||||||||||||||||
Db 3 TCGATCGGGCGGGCGGAGC 22

RESULT 8
AX377574
LOCUS          AX377574      48 bp      DNA      linear      PAT 18-MAR-2002
DEFINITION     Sequence 51 from Patent WO0212553.
ACCESSION      AX377574
VERSION        AX377574.1 GI:19573760
KEYWORDS
SOURCE         synthetic construct.
ORGANISM       synthetic construct
              artificial sequences.
REFERENCE      1
AUTHORS       Kappel,A., Polakowski,T., Pignot,M., Windhab,N., Behrensdoerf,H. and
              Muth,J.
TITLE         Method for detecting mutations in nucleotide sequences
JOURNAL        Patent: WO 0212553-A 51 14-FEB-2002;
              Nanogen
              Recognomics GmbH (DE)
FEATURES       Location/Qualifiers
               1..48
               /organism="synthetic construct"
               /db_xref="taxon:32630"
               /note="Beschreibung der kunstlichen Sequenz:
               Hairpin-Oligonucleotid"
BASE COUNT    6 a      16 c      16 g      10 t
ORIGIN
Query Match   100.0%; Score 20; DB 6; Length 48;
Best Local Similarity 100.0%; Pred. No. 3e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGATCGGGCGGGCGGAGC 20
   ||||||||||||||||||
Db 3 TCGATCGGGCGGGCGGAGC 22

RESULT 9
AX377574/c
LOCUS          AX377574      48 bp      DNA      linear      PAT 18-MAR-2002
DEFINITION     Sequence 51 from Patent WO0212553.
ACCESSION      AX377574
VERSION        AX377574.1 GI:19573760
KEYWORDS
SOURCE         synthetic construct.
ORGANISM       synthetic construct
              artificial sequences.
REFERENCE      1
AUTHORS       Kappel,A., Polakowski,T., Pignot,M., Windhab,N., Behrensdoerf,H. and
              Muth,J.
TITLE         Method for detecting mutations in nucleotide sequences
JOURNAL        Patent: WO 0212553-A 51 14-FEB-2002;
              Nanogen
              Recognomics GmbH (DE)
FEATURES       Location/Qualifiers
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Best Local Similarity 100.0%; Pred. No. 3e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGATCGGGCGGGCGGAGC 20
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Db 46 TCGATCGGGCGGGCGGAGC 27

RESULT 10
AX127457
LOCUS          AX127457      49 bp      DNA      linear      PAT 15-MAY-2001
DEFINITION     Sequence 1 from Patent WO0131057.
ACCESSION      AX127457
VERSION        AX127457.1 GI:14134020
KEYWORDS
SOURCE         synthetic construct.
ORGANISM       synthetic construct
              artificial sequences.
REFERENCE      1 (bases 1 to 49)
AUTHORS       Muth,J. and Windhab,N.
TITLE         Double-strand nucleic acid probes and the use thereof
JOURNAL        Patent: WO 0131057-A 1 03-MAY-2001;
              Aventis
              Research & Technologies GmbH & Co KG (DE)
FEATURES       Location/Qualifiers
               1..49

```

```

Muth,J.
TITLE         Method for detecting mutations in nucleotide sequences
JOURNAL        Patent: WO 0212553-A 51 14-FEB-2002;
              Nanogen
              Recognomics GmbH (DE)
FEATURES       Location/Qualifiers
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               /note="Beschreibung der kunstlichen Sequenz:
               Hairpin-Oligonucleotid"
BASE COUNT    6 a      16 c      16 g      10 t
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Best Local Similarity 100.0%; Pred. No. 3e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGATCGGGCGGGCGGAGC 20
   ||||||||||||||||||
Db 3 TCGATCGGGCGGGCGGAGC 22

RESULT 9
AX377574/c
LOCUS          AX377574      48 bp      DNA      linear      PAT 18-MAR-2002
DEFINITION     Sequence 51 from Patent WO0212553.
ACCESSION      AX377574
VERSION        AX377574.1 GI:19573760
KEYWORDS
SOURCE         synthetic construct.
ORGANISM       synthetic construct
              artificial sequences.
REFERENCE      1
AUTHORS       Kappel,A., Polakowski,T., Pignot,M., Windhab,N., Behrensdoerf,H. and
              Muth,J.
TITLE         Method for detecting mutations in nucleotide sequences
JOURNAL        Patent: WO 0212553-A 51 14-FEB-2002;
              Nanogen
              Recognomics GmbH (DE)
FEATURES       Location/Qualifiers
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DEFINITION     Sequence 1 from Patent WO0131057.
ACCESSION      AX127457
VERSION        AX127457.1 GI:14134020
KEYWORDS
SOURCE         synthetic construct.
ORGANISM       synthetic construct
              artificial sequences.
REFERENCE      1 (bases 1 to 49)
AUTHORS       Muth,J. and Windhab,N.
TITLE         Double-strand nucleic acid probes and the use thereof
JOURNAL        Patent: WO 0131057-A 1 03-MAY-2001;
              Aventis
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RESULT 11
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LOCUS AX127457 49 bp DNA linear PAT 15-MAY-2001
DEFINITION Sequence 1 from Patent WO0131057.
ACCESSION AX127457
VERSION AX127457.1 GI:14134020
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequences.
REFERENCE 1 (bases 1 to 49)
AUTHORS Muth,J. and Windhab,N.
TITLE Double-strand nucleic acid probes and the use thereof
JOURNAL Patent: WO 0131057-A 1 03-MAY-2001;
Aventis Research & Technologies GmbH & Co KG (DE)
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Location/Qualifiers
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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RESULT 12
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LOCUS AX127459 49 bp DNA linear PAT 15-MAY-2001
DEFINITION Sequence 3 from Patent WO0131057.
ACCESSION AX127459
VERSION AX127459.1 GI:14134028
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequences.
REFERENCE 1 (bases 1 to 49)
AUTHORS Muth,J. and Windhab,N.
TITLE Double-strand nucleic acid probes and the use thereof
JOURNAL Patent: WO 0131057-A 3 03-MAY-2001;
Aventis Research & Technologies GmbH & Co KG (DE)
FEATURES
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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LOCUS AX127459 49 bp DNA linear PAT 15-MAY-2001
DEFINITION Sequence 3 from Patent WO0131057.
ACCESSION AX127459
VERSION AX127459.1 GI:14134028
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequences.
REFERENCE 1 (bases 1 to 49)
AUTHORS Muth,J. and Windhab,N.
TITLE Double-strand nucleic acid probes and the use thereof
JOURNAL Patent: WO 0131057-A 3 03-MAY-2001;
Aventis Research & Technologies GmbH & Co KG (DE)
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 47 TCGATCGGGCGGGCGGAGC 28

RESULT 14
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LOCUS AX377573 49 bp DNA linear PAT 18-MAR-2002
DEFINITION Sequence 50 from Patent WO0212553.
ACCESSION AX377573
VERSION AX377573.1 GI:19573759
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequences.
REFERENCE 1
AUTHORS Kappel,A., Polakowski,T., Pignot,M., Windhab,N., Behrensdoerf,H. and
Muth,J.
TITLE Method for detecting mutations in nucleotide sequences
JOURNAL Patent: WO 0212553-A 50 14-FEB-2002;
Nanogen Recognomics GmbH (DE)
FEATURES
Location/Qualifiers
source
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/organism="synthetic construct"
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Hairpin-Oligonucleotid"
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Qy 1 TCGATCGGGCGGGCGGAGC 20
Db 47 TCGATCGGGCGGGCGGAGC 28
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DEFINITION Sequence 825 from Patent WO0122972.
ACCESSION  AX104633
VERSION    AX104633.1  GI:13920830
KEYWORDS
SOURCE     synthetic construct.
           synthetic construct.
           artificial sequences.
ORGANISM
REFERENCE  1 (bases 1 to 21)
AUTHORS   Krieg,A.M., Schetter,C. and Vollmer,J.C.
TITLE     Immunostimulatory nucleic acids
JOURNAL   Patent: WO 0122972-A 825 05-APR-2001;
          UNIVERSITY OF IOWA RESEARCH FOUNDATION (US) ; Coley Pharmaceutical
          GmbH (DE)
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Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  TCGATCGGGCGGGCGGAG 19
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Db      3  TCGATCGGGCGGGCGGAG 21

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GenCore version 5.1.3
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OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:05:40 ; Search time 22.2464 Seconds
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Title: US-09-355-254F-11

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Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 153338381 residues

Total number of hits satisfying chosen parameters: 687286

Minimum DB seq length: 0
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Post-processing: Minimum Match 0%
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Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	20	100.0	21	2	US-08-507-750-2
3	20	100.0	21	3	US-08-764-522A-2
4	20	100.0	21	3	US-08-764-528-2
5	20	100.0	21	3	US-08-872-859-2
6	18.4	92.0	21	1	US-08-283-591-15
7	18.4	92.0	21	1	US-08-210-880B-3
8	18.4	92.0	21	2	US-08-632-275-1
9	18.4	92.0	21	3	US-08-771-411-3
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19	12.8	64.0	22	1	US-08-339-152A-7
20	12.8	64.0	27	4	US-09-564-805-204
21	12.8	64.0	44	1	US-08-458-084-17
22	12.8	64.0	44	1	US-08-205-508-17
23	12.8	64.0	44	5	PCT-US95-02945-17
24	12.6	63.0	20	4	US-09-326-186B-171
25	12.6	63.0	30	3	US-08-913-842-52
26	12.6	63.0	38	4	US-08-341-560B-18
27	12.6	63.0	38	4	US-09-139-802-207

28	12.6	63.0	38	4	US-09-425-638A-3	Sequence 3, Appli
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33	12.2	61.0	20	1	US-08-129-719-13	Sequence 13, Appl
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35	12.2	61.0	20	1	US-08-569-959-13	Sequence 13, Appl
36	12.2	61.0	20	2	US-08-229-528-37	Sequence 37, Appl
37	12.2	61.0	20	3	US-09-429-323-77	Sequence 37, Appl
38	12.2	61.0	35	1	US-08-310-356-31	Sequence 31, Appl
39	12.2	61.0	35	1	US-08-310-356-35	Sequence 35, Appl
40	12.2	61.0	44	4	US-09-110-959A-11	Sequence 11, Appl
41	12.2	61.0	48	4	US-09-025-769B-94	Sequence 94, Appl
42	12	60.0	19	2	US-08-851-135-1	Sequence 1, Appli
43	12	60.0	34	2	US-08-577-492-16	Sequence 16, Appl
44	12	60.0	34	2	US-09-079-630-16	Sequence 16, Appl
45	12	60.0	66	2	US-08-652-816A-39	Sequence 39, Appl

ALIGNMENTS

RESULT 1
US-08-507-598-2
; Sequence 2, Application US/08507598
; Patent No. 5834188
; GENERAL INFORMATION:
; APPLICANT: HARADA, SHUN-ICHI
; APPLICANT: SAMPATH, T. K.
; APPLICANT: RODAN, GIDEON A.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING
; TITLE OF INVENTION: MORPHOGEN ANALOGS
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PATENT ADMINISTRATOR, TESTA, HURWITZ &
; ADDRESS: THIBEAULT
; STREET: 53 STATE STREET
; CITY: BOSTON
; STATE: MA
; COUNTRY: USA
; ZIP: 02109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/507,598
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: PITCHER, EDMUND R.
; REGISTRATION NUMBER: 27,829
; REFERENCE/DOCKET NUMBER: CRP-107
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617)-248-7000
; TELEFAX: (617)-248-7100
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
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; NAME/KEY: misc_feature
; LOCATION: 1..21
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US-08-507-598-2
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Best Local Similarity 100.0%; Pred. No. 0.058;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 2 GCTTGATGACTCAGCGGAA 21

RESULT 2

US-08-507-750-2
; Sequence 2, Application US/08507750
; Patent No. 5932716
; GENERAL INFORMATION:
; APPLICANT: SAMPATH, T. K.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING
; MORPHOGEN ANALOGS
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PATENT ADMINISTRATOR, TESTA, HURWITZ &
; ADDRESSEE: THIBEAULT
; STREET: 53 STATE STREET
; CITY: BOSTON
; STATE: MA
; COUNTRY: USA
; ZIP: 02109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/507.750
; FILING DATE:
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: FITCHER, EDMUND R.
; REGISTRATION NUMBER: 27,829
; REFERENCE/DOCKET NUMBER: CRP-116
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617)-248-7000
; TELEFAX: (617)-248-7100
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
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US-08-507-750-2
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Best Local Similarity 100.0%; Pred. No. 0.058;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Db 2 GCTTGATGACTCAGCGGAA 21

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US-08-764-522A-2
; Sequence 2, Application US/08764522A
; Patent No. 6090544
; GENERAL INFORMATION:
; APPLICANT: HARADA, SHUN-ICHI
; APPLICANT: SAMPATH, T. K.
; APPLICANT: RODAN, GIDEON A.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING
; MORPHOGEN ANALOGS
; NUMBER OF SEQUENCES: 10
; ...

; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PATENT ADMINISTRATOR, CREATIVE BIOMOLECULES
; STREET: 45 SOUTH STREET
; CITY: HOPKINTON
; STATE: MA
; COUNTRY: USA
; ZIP: 01748
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/764.522A
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: VITO, CHRISTINE C.
; REGISTRATION NUMBER: 39,061
; REFERENCE/DOCKET NUMBER: CRP-126
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617)-248-7000
; TELEFAX: (617)-248-7100
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
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US-08-764-522A-2

Query Match 100.0%; Score 20; DB 3; Length 21;
Best Local Similarity 100.0%; Pred. No. 0.058;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 2 GCTTGATGACTCAGCGGAA 21

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US-08-764-528-2
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; Patent No. 6103491
; GENERAL INFORMATION:
; APPLICANT: SAMPATH, K. T.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING
; MORPHOGEN ANALOGS
; NUMBER OF SEQUENCES: 8
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PATENT ADMINISTRATOR, CREATIVE BIOMOLECULES
; STREET: 45 SOUTH STREET
; CITY: HOPKINTON
; STATE: MA
; COUNTRY: USA
; ZIP: 01748
; COMPUTER READABLE FORM:
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; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/764.528
; FILING DATE:
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: VITO, CHRISTINE C.
; REGISTRATION NUMBER: 39,061

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; REFERENCE/DOCKET NUMBER: CRP-127
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617)-248-7000
; TELEFAX: (617)-248-7100
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
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US-08-764-528-2

Query Match 100.0%; Score 20; DB 3; Length 21;
Best Local Similarity 100.0%; Pred. No. 0.058;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 2 GCTTGATGACTCAGCGGAA 21

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; Patent No. 6110460
; GENERAL INFORMATION:
; APPLICANT: SAMPATH, T. K.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING
; TITLE OF INVENTION: MORPHOGEN ANALOGS
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PATENT ADMINISTRATOR, TESTA, HURWITZ &
; ADDRESSEE: THIBEAULT
; STREET: 53 STATE STREET
; CITY: BOSTON
; STATE: MA
; COUNTRY: USA
; ZIP: 02109
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; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/872,859
; FILING DATE: 11-JUN-1997
; STRANDEDNESS: single
; TOPOLOGY: linear
; NAME: PITCHER, EDMUND R.
; REGISTRATION NUMBER: 27,829
; REFERENCE/DOCKET NUMBER: CRP-116
; TELEPHONE: (617)-248-7000
; TELEFAX: (617)-248-7100
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; TYPE: nucleic acid
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; TOPOLOGY: linear
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; NAME/KEY: misc_feature
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; OTHER INFORMATION: /product= "AP1 SEQUENCE"

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US-08-872-859-2

Query Match 100.0%; Score 20; DB 3; Length 21;
Best Local Similarity 100.0%; Pred. No. 0.058;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 2 GCTTGATGACTCAGCGGAA 21

RESULT 6
US-08-283-591-15
; Sequence 15, Application US/08283591
; Patent No. 5629152
; GENERAL INFORMATION:
; APPLICANT: Ravikumar, Vasulunga
; TITLE OF INVENTION: NOVEL TRISUBSTITUTED -LACTAMS AND
; TITLE OF INVENTION: OLIGO -LACTAMAMIDES
; NUMBER OF SEQUENCES: 28
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Woodcock Washburn Kurtz Mackiewicz and
; ADDRESSEE: No. 5629152rls
; STREET: One Liberty Place - 46th Floor
; CITY: Philadelphia
; STATE: PA
; COUNTRY: U.S.A.
; ZIP: 19103
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
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; APPLICATION NUMBER: US/08/283,591
; FILING DATE: N/A
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME:
; REGISTRATION NUMBER:
; REFERENCE/DOCKET NUMBER:
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 215-568-3100
; TELEFAX: 215-568-3439
; INFORMATION FOR SEQ ID NO: 15:
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; STRANDEDNESS: single
; TOPOLOGY: unknown
US-08-283-591-15

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Best Local Similarity 95.0%; Pred. No. 0.42;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 2 GCTTGATGACTCAGCGGAA 21

RESULT 7
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; Sequence 3, Application US/08210880B
; Patent No. 5641486
; GENERAL INFORMATION:
; APPLICANT: HINRICHS, STEVEN H.
; APPLICANT: ORTEN, DANA J.
; TITLE OF INVENTION: METHOD AND COMPOSITION FOR INHIBITING
; TITLE OF INVENTION: CREB/ATF1 FAMILY DETERMINED TRANSCRIPTION

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; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HENDERSON & STURM
; STREET: 1125 S. 103RD ST., #330
; CITY: OMAHA
; STATE: NE
; COUNTRY: US
; ZIP: 68124
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/210.880B
; FILING DATE: 18-MAR-1994
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: JONDLE, ROBERT J.
; REGISTRATION NUMBER: 33,915
; REFERENCE/DOCKET NUMBER: 63066
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 402-398-9000
; TELEFAX: 402-398-9005
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: unknown
; MOLECULE TYPE: DNA (genomic)
US-08-210-880B-3

Query Match 92.0%; Score 18.4; DB 1; Length 21;
Best Local Similarity 95.0%; Pred. No. 0.42;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GCTTGATGACTCAGCGGAA 20
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Db 2 GCTTGATGACTCAGCGGAA 21
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RESULT 8
US-08-632-275-1/c
; Sequence 1, Application US/08632275
; Patent No. 5840277
; GENERAL INFORMATION:
; APPLICANT: Ghio, Andrew J.
; ADDRESSEE: Bell, Seltzer, Park & Gibson
; STREET: 1211 East Morehead Street
; CITY: Charlotte
; STATE: No. 5840277th Carolina
; COUNTRY: USA
; ZIP: 28234
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/632.275
; FILING DATE: 15-APR-1996
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/413.699
; FILING DATE: 30-MAR-1995
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:

```

```

; NAME: Lipscomb, Ernest B.
; REGISTRATION NUMBER: 24,733
; REFERENCE/DOCKET NUMBER: 8751-5-1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 704-331-6000
; TELEFAX: 704-334-2014
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; FRAGMENT TYPE: linear
US-08-632-275-1

Query Match 92.0%; Score 18.4; DB 2; Length 21;
Best Local Similarity 95.0%; Pred. No. 0.42;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GCTTGATGACTCAGCGGAA 20
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Db 20 GCTTGATGACTCAGCGGAA 1
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RESULT 9
US-08-771-411-3
; Sequence 3, Application US/08771411
; Patent No. 5844096
; GENERAL INFORMATION:
; APPLICANT: HINRICHS, STEVEN H.
; APPLICANT: ORTEN, DANA J.
; TITLE OF INVENTION: METHOD AND COMPOSITION FOR INHIBITING
; CREB/ATF1 FAMILY DETERMINED TRANSCRIPTION
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HENDERSON & STURM
; STREET: 1125 S. 103RD ST., #330
; CITY: OMAHA
; STATE: NE
; COUNTRY: US
; ZIP: 68124
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/771.411
; FILING DATE: 20-DEC-1996
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/210.880
; FILING DATE: 18-MAR-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: JONDLE, ROBERT J.
; REGISTRATION NUMBER: 33,915
; REFERENCE/DOCKET NUMBER: 63066
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 402-398-9000
; TELEFAX: 402-398-9005
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: unknown
; MOLECULE TYPE: DNA (genomic)
US-08-771-411-3

Query Match 92.0%; Score 18.4; DB 2; Length 21;
Best Local Similarity 95.0%; Pred. No. 0.42;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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QY 1 GCTTGATGACTCAGCGGAA 20
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Db 2 GCTTGATGACTCAGCGGAA 21

RESULT 10

US-09-097-929-1/c
; Sequence 1, Application US/09097929
; Patent No. 6024940
; GENERAL INFORMATION:
; APPLICANT: Ghio, Andrew J.
; APPLICANT: Kennedy, Thomas P.
; TITLE OF INVENTION: Treatment of Chronic Pulmonary
; TITLE OF INVENTION: Inflammation
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Bell, Seltzer, Park & Gibson
; STREET: 1211 East Morehead Street
; CITY: Charlotte
; STATE: No. 6024940th Carolina
; COUNTRY: USA
; ZIP: 28234
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/097,929
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; PRIOR APPLICATION NUMBER: 08/632,275
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Lipscomb, Ernest B.
; REGISTRATION NUMBER: 24,733
; REFERENCE/DOCKET NUMBER: 8751-5-1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 704-331-6000
; TELEFAX: 704-334-2014
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; FRAGMENT TYPE: linear
US-09-097-929-1

Query Match 92.0%; Score 18.4; DB 3; Length 21;
Best Local Similarity 95.0%; Pred. No. 0.42; Mismatches 0; Gaps 0;
Matches 19; Conservative 0; Indels 1; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20
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Db 20 GCTTGATGACTCAGCGGAA 1

RESULT 11

US-09-021-247-8
; Sequence 8, Application US/09021247
; Patent No. 6225444
; GENERAL INFORMATION:
; APPLICANT: Shashoua, Victor E.
; TITLE OF INVENTION: NEUROPROTECTIVE PEPTIDES AND USES THEREOF
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Wolf, Greenfield & Sacks, P.C.
; STREET: 600 Atlantic Avenue
; CITY: Boston
; STATE: MA

; COUNTRY: USA
; ZIP: 02210
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/021,247
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Van Amsterdam, John R.
; REGISTRATION NUMBER: 40,212
; REFERENCE/DOCKET NUMBER: N0260/7023
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 617-720-3500
; TELEFAX: 617-720-2441
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 nucleotides
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: oligonucleotide
; HYPOTHETICAL: NO
US-09-021-247-8

Query Match 92.0%; Score 18.4; DB 4; Length 21;
Best Local Similarity 95.0%; Pred. No. 0.42; Mismatches 19; Conservative 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20
||||||| |||||||
Db 2 GCTTGATGACTCAGCGGAA 21

RESULT 12

US-08-088-661F-8
; Sequence 8, Application US/08088661F
; Patent No. 6228982
; GENERAL INFORMATION:
; APPLICANT: No. 6228982den, Bengt
; APPLICANT: Wittung, Pernilla
; APPLICANT: Buchardt, Ole
; APPLICANT: Egholm, Michael
; APPLICANT: Nielsen, Peter E.
; APPLICANT: Berg, Rolf
; TITLE OF INVENTION: Double-Stranded Peptide Nucleic Acids
; FILE REFERENCE: ISIS1108
; CURRENT APPLICATION NUMBER: US/08/088,661F
; CURRENT FILING DATE: 1993-07-02
; PRIOR APPLICATION NUMBER: 08/054,363
; PRIOR FILING DATE: 1993-04-26
; PRIOR APPLICATION NUMBER: PCT/EP92/01219
; PRIOR FILING DATE: 1992-05-19
; NUMBER OF SEQ ID NOS: 42
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 8
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: No. 6228982el Sequence
US-08-088-661F-8

Query Match 92.0%; Score 18.4; DB 4; Length 21;
Best Local Similarity 95.0%; Pred. No. 0.42; Mismatches 19; Conservative 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20
||||||| |||||||
Db 2 GCTTGATGACTCAGCGGAA 21

RESULT 13
US-08-088-661F-42
; Sequence 42, Application US/08088661F
; Patent No. 6228982
; GENERAL INFORMATION:
; APPLICANT: No. 6228982den, Benget
; APPLICANT: Wittung, Pernilla
; APPLICANT: Buchardt, Ole
; APPLICANT: Egholm, Michael
; APPLICANT: Nielsen, Peter E.
; APPLICANT: Berg, Rolf
; TITLE OF INVENTION: Double-Stranded Peptide Nucleic Acids
; FILE REFERENCE: IS11108
; CURRENT APPLICATION NUMBER: US/08/088,661F
; CURRENT FILING DATE: 1993-07-02
; PRIOR APPLICATION NUMBER: 08/054,363
; PRIOR FILING DATE: 1993-04-26
; PRIOR APPLICATION NUMBER: PCT/EP92/01219
; PRIOR FILING DATE: 1992-05-19
; NUMBER OF SEQ ID NOS: 42
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 42
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: No. 6228982el Sequence
US-08-088-661F-42

Query Match 92.0%; Score 18.4; DB 4; Length 21;
Best Local Similarity 95.0%; Pred. No. 0.42;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20
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Db 2 GCTTGGTGACTCAGCGGAA 21

RESULT 14
US-08-203-198-1
; Sequence 1, Application US/08203198
; Patent No. 5512462
; GENERAL INFORMATION:
; APPLICANT: Cheng, Suzanne
; TITLE OF INVENTION: Methods and Reagents for the Polymerase
; NUMBER OF SEQUENCES: 32
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Hoffmann-La Roche Inc.
; STREET: 340 Kingsland Street
; CITY: Nutley
; STATE: NJ
; COUNTRY: U.S.A.
; ZIP: 07110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/203,198
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Petry Ph.D., Douglas A.
; REGISTRATION NUMBER: 35,321
; REFERENCE/DOCKET NUMBER: 8894
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (510) 814-2974
; TELEFAX: (510) 814-2977
; INFORMATION FOR SEQ ID NO: 1:

; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-08-203-198-1
Query Match 69.0%; Score 13.8; DB 1; Length 21;
Best Local Similarity 88.2%; Pred. No. 1.2e+02;
Matches 15; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 GCTTGATGACTCAGCG 17
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Db 4 GCTTTATGACTCTGCG 20
RESULT 15
US-07-791-213D-68/C
; Sequence 68, Application US/07791213D
; Patent No. 5409895
; GENERAL INFORMATION:
; APPLICANT: MORISHITA, Hideaki
; APPLICANT: KANAMORI, Toshinori
; APPLICANT: NOBUHARA, Masahiro
; TITLE OF INVENTION: POLYPEPTIDE, DNA FRAGMENT ENCODING THE
; TITLE OF INVENTION: SAME AND PROCESS FOR PRODUCING THE SAME, AND ENZYME
; TITLE OF INVENTION: INHIBITION PROCESS, DRUG COMPOSITION AND METHODS OF
; TITLE OF INVENTION: TREATING USING THE SAME
; NUMBER OF SEQUENCES: 108
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Burns, Doane, Swecker & Mathis
; STREET: P.O. Box 1404
; CITY: Alexandria
; STATE: Virginia
; COUNTRY: United States
; ZIP: 22313-1404
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/791,213D
; FILING DATE: 13-NOV-1991
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: JP 2-306745
; FILING DATE: 13-NOV-1990
; ATTORNEY/AGENT INFORMATION:
; NAME: Meuth, Donna M
; REGISTRATION NUMBER: 36,607
; REFERENCE/DOCKET NUMBER: 029650-032
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703) 836-6620
; TELEFAX: (703) 836-2021
; INFORMATION FOR SEQ ID NO: 68:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 26 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: complement (1..26)
US-07-791-213D-68
Query Match 68.0%; Score 13.6; DB 1; Length 26;
Best Local Similarity 80.0%; Pred. No. 1.6e+02;
Matches 16; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
QY 1 GCTTGATGACTCAGCGGAA 20

Db ||| ||| ||| ||| ||| ||| |||
 26 GCTGGATCCCTCAGCCGAAA 7

Search completed: December 12, 2002, 01:41:48
Job time : 24.2464 secs

GenCore version 5.1.3
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:06:16 ; Search time 318.116 Seconds
(without alignments)
1829.698 Million cell updates/sec

Title: US-09-355-254F-14

Perfect score: 20
Sequence: 1 agcggggcgagcgggggcg 20

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0
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Post-processing: Minimum Match 0%
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Listing first 45 summaries

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- 11: gb_sts.*
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- 40: em_htgo_mus.*
- 41: em_htgo_other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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1	20	100.0	20	6	A89793	A89793 Sequence 15
2	20	100.0	20	6	A90880	A90880 Sequence 15
3	20	100.0	20	6	AX023403	AX023403 Sequence
4	20	100.0	20	6	AX455554	AX455554 Sequence
5	20	100.0	30	6	AX472520	AX472520 Sequence
6	20	100.0	30	6	AX476834	AX476834 Sequence
7	20	100.0	30	6	AX476857	AX476857 Sequence
8	16.4	82.0	30	6	AR063474	AR063474 Sequence
9	16.4	82.0	30	6	I28154	I28154 Sequence 11
10	14.2	71.0	47	6	AX128389	AX128389 Sequence
11	14.2	71.0	50	6	AR032813	AR032813 Sequence
12	14.2	71.0	50	6	AR209477	AR209477 Sequence
13	14.2	71.0	50	6	I29553	I29553 Sequence 42
14	14.2	71.0	50	6	I91227	I91227 Sequence 42
15	14.2	71.0	51	6	AX158283	AX158283 Sequence
16	13.8	69.0	72	9	HSNDSADAE	Z22319 H.sapiens D
17	13.8	69.0	100	11	AF235063	AF235063 Mus muscu
18	13.6	68.0	30	6	AX472521	AX472521 Sequence
19	13.6	68.0	30	6	AX476835	AX476835 Sequence
20	13.6	68.0	40	6	I86249	I86249 Sequence 3
21	13.6	68.0	44	6	AX157237	AX157237 Sequence
22	13.6	68.0	50	10	MMU41966	U41966 Mus musculu
23	13.6	68.0	51	6	A42081	A42081 Sequence 24
24	13.6	68.0	98	6	AR017637	AR017637 Sequence
25	13.6	68.0	98	6	AR094814	AR094814 Sequence
26	13.6	68.0	98	6	AR165473	AR165473 Sequence
27	13.6	68.0	99	8	AY033466	AY033466 zea mays
28	13.6	68.0	100	6	I28243	I28243 Sequence 1
29	13.4	67.0	24	6	AR058202	AR058202 Sequence
30	13.4	67.0	24	6	AR152031	AR152031 Sequence
31	13.4	67.0	24	6	I50793	I50793 Sequence 24
32	13.4	67.0	30	6	AX167104	AX167104 Sequence
33	13.4	67.0	33	6	AX202261	AX202261 Sequence
34	13.2	66.0	19	6	AX129905	AX129905 Sequence
35	13.2	66.0	19	6	AX129926	AX129926 Sequence
36	13.2	66.0	19	6	AX202627	AX202627 Sequence
37	13.2	66.0	21	6	A88115	A88115 Sequence 26
38	13.2	66.0	21	6	A90082	A90082 Sequence 26
39	13.2	66.0	22	6	A33211	A33211 Synthetic H
40	13.2	66.0	47	6	AR154447	AR154447 Sequence
41	13.2	66.0	50	6	AX159936	AX159936 Sequence
42	13.2	66.0	51	6	AX159935	AX159935 Sequence
43	13.2	66.0	54	6	AR032427	AR032427 Sequence
44	13.2	66.0	54	6	AR032648	AR032648 Sequence
45	13.2	66.0	54	6	AR209091	AR209091 Sequence

ALIGNMENTS

RESULT 1

A89793

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

A89793
Sequence 15 from Patent WO9832462.
A89793
A89793.1 GI:6738307

20 bp

DNA

linear

PAT 22-JAN-2000

unidentified.
unidentified.
unclassified.

1 (bases 1 to 20)

Lipford, G. B. and Heeg, K.

PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND

OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION

Patent: WO 9832462-A 15 30-JUL-1998;

LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)

FEATURES
source

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/db_xref="taxon:32644" 0 t

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ORIGIN

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Best Local Similarity 100.0%; Pred. No. 2.2e+03;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCGGGGCGAGCGGGGCG 20

Db 1 AGCGGGGCGAGCGGGGCG 20

RESULT 2

A90880

LOCUS A90880 20 bp DNA linear PAT 22-JAN-2000

DEFINITION Sequence 15 from Patent EP0855184.

ACCESSION A90880

VERSION A90880.1 GI:6739295

KEYWORDS unidentified.

SOURCE unidentified.

ORGANISM unclassified.

REFERENCE 1 (bases 1 to 20)

AUTHORS Heeg,K.P. and Lipford,G.B.

TITLE Pharmaceutical composition comprising a polynucleotide and an

antigen especially for vaccination

JOURNAL Patent: EP 0855184-A 15 29-JUL-1998;

HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)

FEATURES

source

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BASE COUNT 2 a 4 c 14 g 0 t

ORIGIN

Query Match 100.0%; Score 20; DB 6; Length 20;

Best Local Similarity 100.0%; Pred. No. 2.2e+03;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCGGGGCGAGCGGGGCG 20

Db 1 AGCGGGGCGAGCGGGGCG 20

RESULT 3

AX023403

LOCUS AX023403 20 bp DNA linear PAT 15-SEP-2000

DEFINITION Sequence 18 from Patent WO0014217.

ACCESSION AX023403

VERSION AX023403.1 GI:10183803

KEYWORDS synthetic construct.

SOURCE synthetic construct.

ORGANISM artificial sequences.

REFERENCE 1 (bases 1 to 20)

AUTHORS Lipford,G.B., Heeg,K. and Wagner,H.

TITLE G-motif oligonucleotides and uses thereof

JOURNAL Patent: WO 0014217-A 18 16-MAR-2000;

LIPFORD GRAYSON B (DE); HEEG KLAUS (DE); WAGNER HERMANN (DE);

CPG IMMUNOPHARMACEUTICALS GMBH (DE)

FEATURES

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/organism="synthetic construct"

/db_xref="taxon:32630"

/note="synthetic, no natural origin"

BASE COUNT 2 a 4 c 14 g 0 t

ORIGIN

Query Match 100.0%;

Best Local Similarity 100.0%; Score 20; DB 6; Length 20;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCGGGGCGAGCGGGGCG 20

Db 1 AGCGGGGCGAGCGGGGCG 20

RESULT 4

AX455554

LOCUS AX455554

DEFINITION Sequence 31 from Patent WO0222809.

ACCESSION AX455554

VERSION AX455554.1 GI:21714622

KEYWORDS synthetic construct.

SOURCE synthetic construct.

ORGANISM artificial sequences.

REFERENCE 1

AUTHORS Bauer,S., Lipford,G. and Wagner,H.

TITLE Process for high throughput screening of cpg-based

immuno-agonist/antagonist

JOURNAL Patent: WO 0222809-A 31 21-MAR-2002;

Coley Pharmaceutical GmbH (DE)

FEATURES Location/Qualifiers

source

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/organism="synthetic construct"

/db_xref="taxon:32630"

/note="Synthetic oligonucleotide"

BASE COUNT 2 a 4 c 14 g 0 t

ORIGIN

Query Match 100.0%; Score 20; DB 6; Length 20;

Best Local Similarity 100.0%; Pred. No. 2.2e+03;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCGGGGCGAGCGGGGCG 20

Db 1 AGCGGGGCGAGCGGGGCG 20

RESULT 5

AX472520

LOCUS AX472520

DEFINITION Sequence 15 from Patent WO02052039.

ACCESSION AX472520

VERSION AX472520.1 GI:22207424

KEYWORDS synthetic construct.

SOURCE synthetic construct.

ORGANISM artificial sequences.

REFERENCE 1

AUTHORS Blais,Y., Rousseau,P., Leblanc,B. and Camato,R.N.

TITLE Methods for selecting and producing selective pharmaceutical

compounds and compositions using an established genetically altered

cell-based library responsive to transcription factors; genetic

constructs and library therefor

JOURNAL Patent: WO 02052039-A 15 04-JUL-2002;

Geneka Biotechnology Inc. (CA)

FEATURES Location/Qualifiers

source

1..30

/organism="synthetic construct"

/db_xref="taxon:32630"

/note="oligonucleotide"

BASE COUNT 5 a 7 c 17 g 1 t

ORIGIN

Query Match 100.0%; Score 20; DB 6; Length 30;

Best Local Similarity 100.0%; Pred. No. 2e+03;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCGGGGCGAGCGGGGCG 20

```
Db 7 AGCGGGGGCGAGCGGGGCG 26
|||||
AX476834
LOCUS AX476834 30 bp DNA linear PAT 12-AUG-2002
DEFINITION Sequence 11 from Patent WO02052037.
ACCESSION AX476834
VERSION AX476834.1 GI:22216110
KEYWORDS synthetic construct.
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE 1
AUTHORS Larose,A.M., Rousseau,P., Leblanc,B. and Camato,R.
TITLE Method for screening and/or identifying factors that bind to
JOURNAL nucleic acids
PATENT: WO 02052037-A 11 04-JUL-2002;
Geneka Biotechnology Inc. (CA)
FEATURES
source
Location/Qualifiers
1..30
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="NABE-probes"
BASE COUNT 5 a 7 c 17 g 1 t
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Query Match 100.0%; Score 20; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGCGGGGGCGAGCGGGGCG 20
|||||
Db 7 AGCGGGGGCGAGCGGGGCG 26
|||||
AX476857
LOCUS AX476857 30 bp DNA linear PAT 12-AUG-2002
DEFINITION Sequence 34 from Patent WO02052037.
ACCESSION AX476857
VERSION AX476857.1 GI:22216133
KEYWORDS synthetic construct.
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE 1
AUTHORS Larose,A.M., Rousseau,P., Leblanc,B. and Camato,R.
TITLE Method for screening and/or identifying factors that bind to
JOURNAL nucleic acids
PATENT: WO 02052037-A 34 04-JUL-2002;
Geneka Biotechnology Inc. (CA)
FEATURES
source
Location/Qualifiers
1..30
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Double stranded NABE"
BASE COUNT 5 a 7 c 17 g 1 t
ORIGIN
Query Match 100.0%; Score 20; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGCGGGGGCGAGCGGGGCG 20
|||||
Db 7 AGCGGGGGCGAGCGGGGCG 26
|||||
AX128389
LOCUS AX128389 47 bp DNA linear PAT 15-MAY-2001
DEFINITION Sequence 50 from Patent WO0130843.
ACCESSION AX128389
VERSION AX128389.1 GI:14134897
KEYWORDS synthetic construct.
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE 1 (bases 1 to 47)
AUTHORS Barbas,C.F., Kadan,M. and Beerli,R.
TITLE Ligand activated transcriptional regulator proteins
JOURNAL Patent: WO 0130843-A 50 03-MAY-2001;
Novartis AG (CH); The Scripps Research Institute (US)
FEATURES
source
Location/Qualifiers
1..47
/organism="synthetic construct"
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/db_xref="taxon:32630"
/note="Recombinant molecule"
BASE COUNT      11 a      11 c      20 g      5 t
ORIGIN

Query Match      71.0%; Score 14.2; DB 6; Length 47;
Best Local Similarity 84.2%; Pred. No. 2.3e+05;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY  2 GCGGGGGCGAGCGGGGCGG 20
    ||||| ||||| |||||
Db   20 GCAGGGCGGAGCGGTGGCG 38

RESULT 11
AR032813/c
LOCUS      AR032813      50 bp      DNA
DEFINITION Sequence 425 from patent US 5869241.
ACCESSION  AR032813
VERSION     AR032813.1 GI:5948418
KEYWORDS    Unknown.
SOURCE      Unknown.
ORGANISM    Unclassified.
REFERENCE   1 (bases 1 to 50)
AUTHORS     Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.
TITLE       Method of determining DNA sequence preference of a DNA-binding
JOURNAL     molecule
FEATURES    Patent: US 5869241-A 425 09-FEB-1999;
            Location/Qualifiers
            source
            1..50
            /organism="unknown"
BASE COUNT      6 a      23 c      17 g      4 t
ORIGIN

Query Match      71.0%; Score 14.2; DB 6; Length 50;
Best Local Similarity 84.2%; Pred. No. 2.3e+05;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY  2 GCGGGGGCGAGCGGGGCGG 20
    ||||| ||||| |||||
Db   23 GCGGGGGCGGCGGGCGGCG 5

RESULT 12
AR209477/c
LOCUS      AR209477      50 bp      DNA
DEFINITION Sequence 425 from patent US 6384208.
ACCESSION  AR209477
VERSION     AR209477.1 GI:21510913
KEYWORDS    Unknown.
SOURCE      Unknown.
ORGANISM    Unclassified.
REFERENCE   1 (bases 1 to 50)
AUTHORS     Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.
TITLE       Sequence directed DNA binding molecules compositions and methods
JOURNAL     Patent: US 6384208-A 425 07-MAY-2002;
FEATURES    Location/Qualifiers
            source
            1..50
            /organism="unknown"
BASE COUNT      6 a      23 c      17 g      4 t
ORIGIN

Query Match      71.0%; Score 14.2; DB 6; Length 50;
Best Local Similarity 84.2%; Pred. No. 2.3e+05;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY  2 GCGGGGGCGAGCGGGGCGG 20
    ||||| ||||| |||||
Db   23 GCGGGGGCGGCGGGCGGCG 5

RESULT 13
I29553/c
LOCUS      I29553      50 bp      DNA
DEFINITION Sequence 425 from patent US 5578444.
ACCESSION  I29553
VERSION     I29553.1 GI:1820344
KEYWORDS    Unknown.
SOURCE      Unknown.
ORGANISM    Unclassified.
REFERENCE   1 (bases 1 to 50)
AUTHORS     Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.
TITLE       Sequence-directed DNA-binding molecules compositions and methods
JOURNAL     Patent: US 5578444-A 425 26-NOV-1996;
FEATURES    Location/Qualifiers
            source
            1..50
            /organism="unknown"
BASE COUNT      6 a      23 c      17 g      4 t
ORIGIN

Query Match      71.0%; Score 14.2; DB 6; Length 50;
Best Local Similarity 84.2%; Pred. No. 2.3e+05;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY  2 GCGGGGGCGAGCGGGGCGG 20
    ||||| ||||| |||||
Db   23 GCGGGGGCGGCGGGCGGCG 5

RESULT 14
I91227/c
LOCUS      I91227      50 bp      DNA
DEFINITION Sequence 425 from patent US 5726014.
ACCESSION  I91227
VERSION     I91227.1 GI:3935697
KEYWORDS    Unknown.
SOURCE      Unknown.
ORGANISM    Unclassified.
REFERENCE   1 (bases 1 to 50)
AUTHORS     Edwards,C.A., Cantor,C.R., Andrews,B.M. and Turin,L.M.
TITLE       Screening assay for the detection of DNA-binding molecules
JOURNAL     Patent: US 5726014-A 425 10-MAR-1998;
FEATURES    Location/Qualifiers
            source
            1..50
            /organism="unknown"
BASE COUNT      6 a      23 c      17 g      4 t
ORIGIN

Query Match      71.0%; Score 14.2; DB 6; Length 50;
Best Local Similarity 84.2%; Pred. No. 2.3e+05;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY  2 GCGGGGGCGAGCGGGGCGG 20
    ||||| ||||| |||||
Db   23 GCGGGGGCGGCGGGCGGCG 5

RESULT 15
AX158283/c
LOCUS      AX158283      51 bp      DNA
DEFINITION Sequence 1611 from Patent WO0140521.
ACCESSION  AX158283
VERSION     AX158283.1 GI:14539614
KEYWORDS    human.
SOURCE      Homo sapiens
ORGANISM    Homo sapiens
REFERENCE   1 (bases 1 to 51)
AUTHORS     Shimkets,R.A. and Leach,M.
TITLE       Nucleic acids containing single nucleotide polymorphisms and
            methods of use thereof
```



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JOURNAL Patent: WO 0140521-A 1611 07-JUN-2001;
FEATURES Curagen Corporation (US)
source Location/Qualifiers
1. .51
/organism="Homo sapiens"
/db_xref="taxon:9606"
misc_feature 26
/notes="1 of 2 allelic variants (1612 is other entry)
Accession number cg32120097"
BASE COUNT 6 a 22 c 15 g 8 t
ORIGIN

Query Match 71.0%; Score 14.2; DB 6; Length 51;
Best Local Similarity 84.2%; Pred. No. 2.2e+05;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGCGGGGGCGAGCGGGGC 19
| | | | | | | | | | | | | | | |
Db 43 AGCGGGGGCCAGCGGGGAGC 25

Search completed: December 12, 2002, 02:55:54
Job time : 323.116 secs

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GenCore version 5.1.3
Copyright (c) 1993 - 2002 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:07:11 ; Search time 815.797 Seconds
(without alignments)
397.047 Million cell updates/sec

Title: US-09-355-254F-8

Perfect score: 20

Sequence: 1 gattgctgacgtcagag 20

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 357874

Minimum DB seq length: 0

Maximum DB seq length: 100

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:**

1: em_estba:*
2: em_esthum:*
3: em_estlin:*
4: em_estnu:*
5: em_estov:*
6: em_estpl:*
7: em_estro:*
8: em_hic:*
9: gb_est1:*
10: gb_est2:*
11: gb_hic:*
12: gb_est3:*
13: gb_est4:*
14: gb_est5:*
15: em_estfun:*
16: em_estom:*
17: gb_gss:*
18: em_gss_hum:*
19: em_gss_inv:*
20: em_gss_pln:*
21: em_gss_vrt:*
22: em_gss_fun:*
23: em_gss_mam:*
24: em_gss_mus:*
25: em_gss_other:*
26: em_gss_pro:*
27: em_gss_rod:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	14.2	71.0	78	17	AZ871578
C 2	13.4	67.0	50	17	AZ492887
C 3	13.2	66.0	49	17	BH861707
C 4	13.2	66.0	50	9	AU107195
C 5	13.2	66.0	67	9	AA721395
C 6	13.2	66.0	71	9	AA702828

C 7	13.2	66.0	91	17	AZ512653
C 8	13.2	66.0	93	9	AA560086
C 9	13.2	66.0	99	12	BF458655
C 10	12.8	64.0	34	17	AQ073796
C 11	12.8	64.0	40	9	AA848120
C 12	12.8	64.0	50	9	AU105994
C 13	12.8	64.0	69	17	BH256492
C 14	12.8	64.0	86	9	AI204909
C 15	12.8	64.0	88	9	AU260302
C 16	12.8	64.0	99	14	F26219
C 17	12.6	63.0	73	10	BE095402
C 18	12.6	63.0	73	17	BH862484
C 19	12.6	63.0	81	9	AA580289
C 20	12.6	63.0	84	10	AV952690
C 21	12.6	63.0	84	17	AZ312968
C 22	12.6	63.0	91	17	AZ494456
C 23	12.6	63.0	95	17	AZ810823
C 24	12.6	63.0	98	17	BH853467
C 25	12.4	62.0	93	14	W31755
C 26	12.4	62.0	94	17	AZ763191
C 27	12.2	61.0	43	17	BH796799
C 28	12.2	61.0	50	9	AU107190
C 29	12.2	61.0	50	9	AU107194
C 30	12.2	61.0	50	9	AU107198
C 31	12.2	61.0	54	17	AZ776635
C 32	12.2	61.0	55	17	AZ920728
C 33	12.2	61.0	64	9	AI253600
C 34	12.2	61.0	64	13	BI598792
C 35	12.2	61.0	74	9	AA218675
C 36	12.2	61.0	79	9	AA600457
C 37	12.2	61.0	80	9	AA985780
C 38	12.2	61.0	82	14	W05256
C 39	12.2	61.0	83	9	AA075933
C 40	12.2	61.0	85	9	AA716566
C 41	12.2	61.0	88	17	AZ430802
C 42	12.2	61.0	90	12	BG400826
C 43	12.2	61.0	97	14	H28506
C 44	12.2	61.0	100	13	BI002964
C 45	12.2	61.0	100	17	AZ780127

ALIGNMENTS

RESULT 1

AZ871578/C

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

AZ871578, 78 bp DNA linear GSS 21-FEB-2001
2M0184G04R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC2M0184G04 R, DNA sequence.

AZ871578

AZ871578.1 GI:13077918

GSS.

house mouse.

Mus musculus

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus;

Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,

Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly

,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.

and Wright,D.,Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts

Unpublished (2000)

Contact: Robert B. Weiss

University of Utah Genome Center

University of Utah

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Class: TDNA tagged.
 Location/Qualifiers
 1. .49
 /organism="Arabidopsis thaliana"
 /strain="Columbia 0"
 /db_xref="taxon:3702"
 /clone="SALK_087868"
 /clone_lib="Arabidopsis thaliana TDNA insertion lines"
 /note="PCR was performed on Arabidopsis thaliana lines each of which contains one or more TDNA insertion elements. The resultant fragment for each line was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at: http://signal.salk.edu/tdna_protocols.html"

BASE COUNT 16 a 14 c 7 g 12 t
 ORIGIN

Query Match 66.0%; Score 13.2; DB 17; Length 49;
 Best Local Similarity 83.3%; Pred. No. 2.8e+04;
 Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 ATTGCCTGACGCTCAGAGA 19
 ||| | ||||| |||||
 Db 36 ATTCTTGACGCTCAGAGA 19

RESULT 4
 AUI07195/c
 LOCUS
 DEFINITION
 50 bp mRNA linear EST 30-AUG-2001
 AUI07195 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
 HEP21410, mRNA sequence.
 AUI07195
 VERSION
 AUI07195.1 GI:13556716
 EST.
 SOURCE
 human.

ORGANISM
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE
 1 (bases 1 to 50)
 AUTHORS
 Suzuki.Y., Taira.H., Tsunoda.T., Mizushima-Sugano.J., Sese,J., Hata
 H., Ota.T., Isogai.T., Tanaka.T., Morishita.S., Okubo.K., Sakaki
 Y., Nakamura.Y., Suyama.A. and Sugano.S.
 TITLE
 Diverse transcriptional initiation revealed by fine, large-scale
 mapping of mRNA start sites
 JOURNAL
 EMBO Rep. 2 (5), 388-393 (2001)
 MEDLINE
 21270072
 COMMENT
 Contact: Yutaka Suzuki
 Department of Virology
 Institute of Medical Science, University of Tokyo
 4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
 Email: ysuzuki@ims.u-tokyo.ac.jp
 Suzuki.Y., Yoshitomo-Nakagawa.K., Maruyama.K., Suyama.A. and Sugano
 S. Construction and characterization of a full length-enriched and
 a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).

FEATURES
 Location/Qualifiers
 1. .50
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="HEP21410"
 /clone_lib="Sugano Homo sapiens cDNA library"
 /note="Differential display comparison of untreated and
 dimethylformate treated U937 cells"
 BASE COUNT 12 a 14 c 14 g 10 t
 ORIGIN

Query Match 66.0%; Score 13.2; DB 9; Length 50;
 Best Local Similarity 83.3%; Pred. No. 2.8e+04;
 Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 TTGCCTGACGCTCAGAG 20
 ||| | ||||| |||||
 Db 41 TTGGCTGACGCTCAACG 24

RESULT 5
 AA721395/c
 LOCUS
 DEFINITION
 67 bp mRNA linear EST 23-JAN-1998
 n273g08.s1 NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:1301150 3',
 mRNA sequence.
 AA721395
 VERSION
 AA721395.1 GI:2737530
 EST.
 SOURCE
 human.
 ORGANISM
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE
 1 (bases 1 to 67)
 AUTHORS
 NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 TITLE
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 UNPUBLISHED (1997)
 JOURNAL
 COMMENT
 Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-r@mail.nih.gov
 Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman,
 Ph.D., Gerald Marti, M.D.
 CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
 Bonaldo, Ph.D.
 CDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
www.bio.llnl.gov/bbrp/image/image.html
 Insert length: 835 Std Error: 0.00
 Seq primer: -40ml3 fwd. ET from Amersham
 High quality sequence stop: 57.

FEATURES
 Location/Qualifiers
 1. 67
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:1301150"
 /clone_lib="NCI_CGAP_GCB1"
 /tissue_type="germinal center B cell"
 /lab_host="DH10B"
 /note="Vector: pT73D-Pac (Pharmacia) with a modified
 polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
 was prepared from human tonsillar cells enriched for
 germinal center B cells by flow sorting (CD20+, IgD-),
 provided by Dr. Louis M. Staudt (NCI), Dr. David Allman
 (NCI) and Dr. Gerald Marti (CBER). CDNA synthesis was
 primed with a Not I - oligo(dT) primer
 [5'-TGTTACCAATCTGAAGTGGAGCGCCCTCATTTTTTTTTTTT-3'
]. Double-stranded cDNA was ligated to Eco RI adaptors
 (Pharmacia), digested with Not I and cloned into the Not I
 and Eco RI sites of the modified pT73 vector. Library
 went through one round of normalization, and was
 constructed by Bento Soares and M. Fatima Bonaldo."
 BASE COUNT 14 a 24 c 15 g 14 t
 ORIGIN

Query Match 66.0%; Score 13.2; DB 9; Length 67;
 Best Local Similarity 83.3%; Pred. No. 3.2e+04;
 Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAG 18
 ||||| ||||| |||||
 Db 30 GATTGCTGACCCACAG 13

RESULT 6
 AA702828/c
 LOCUS
 DEFINITION
 71 bp mRNA linear EST 19-DEC-1997
 z177b06.s1 Soares fetal_liver_spleen_LNFLS_S1 Homo sapiens cDNA
 clone IMAGE:436787 3' similar to contains element CER repetitive
 element ;, mRNA sequence.
 AA702828
 VERSION
 AA702828.1 GI:2705941

small number of additional specialized non-redundant arrays of BMA cDNAs whose availability will be considered under appropriate and limited collaborative arrangements. The following repetitive elements were found in this cDNA sequence: 1-21, >AT-rich#Low_complexity

Seq primer: M13 Forward

POLYA=Yes.

Location/Qualifiers

1. .99

/organism="Mus musculus"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UI-M-B21-blt-g-01-0-UI"

/clone_lib="NIH_BMAP_MHI2_S1"

/dev_stage="27-32 days"

/lab_host="DH10B (Life Technologies)"

/note="Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; The NIH_BMAP_MHI2_S1 library is a subtracted library derived from NIH_BMAP_MHI2. NIH_BMAP_MHI2 is a library derived from mouse hippocampus tissue. For a detailed description of the library from which this clone was derived, please visit our web site at brainest.eng.uiowa.edu."

TAG_LIB="NIH_BMAP_MHI2_S1"

TAG_TISSUE="hippocampus"

TAG_SEQ="TAGCC"

BASE COUNT 18 a 26 c 32 g 23 t

ORIGIN

Query Match 66.0%; Score 13.2; DB 12; Length 99;

Best Local Similarity 83.3%; Pred. No. 3.7e+04;

Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 TTGCTGACGCTCAGAG 20

||||||| |||||||

Db 57 TTGCTGGGTGAGAG 40

RESULT 10

AQ073796/c

LOCUS

DEFINITION

AQ073796 34 bp DNA linear GSS 23-AUG-2000

EP(3)3179 Drosophila melanogaster EP line Drosophila melanogaster genomic sequence recovered from Both 5' and 3' ends of P element, DNA sequence.

ACCESSION

AQ073796.1 GI:3403838

VERSION

GSS.

KEYWORDS

fruit fly.

SOURCE

Drosophila melanogaster

ORGANISM

Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

REFERENCE

Liao,G.-C., Rehm,E.J. and Rubin,G.M.

AUTHORS

Insertion site preferences of the P transposable element in Drosophila melanogaster

TITLE

Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3347-3351 (2000)

JOURNAL

20202638

MEDLINE

Contact: Gerald Rubin

COMMENT

Berkeley Drosophila Genome Project

University of California, Berkeley

LSA Building, Berkeley, CA 94720-3200, USA

Fax: 5106439947

Email: gerry@fruitfly.berkeley.edu

Sequence recovery method was inverse PCR.

Sequence orientation is forward strand relative to 5' end of P element

The P element insertion position is base 15 in the 34 bases. This insertion position refers to the first base of the 8 base target recognition sequence.

Class: transposon-tagged.

FEATURES
source

Location/Qualifiers
1. .34
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/clone_lib="Drosophila melanogaster EP line"
/note="Inverse PCR was performed on Drosophila melanogaster strains each of which contains a single EP transposable element insertion. (The generation of these insertion strains is described in Rorth P, Szabo K, Bailey A, Lavery T, Rehm J, Rubin GM, Weigmann K, Milan M, Benes V, Ansorge W, Cohen SM. 1998. Systematic gain-of-function genetics in Drosophila. Development 6:1049-1057.) The resultant fragment for each strain was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at http://fruitfly.berkeley.edu/p_disrupt/inverse_pcr.html."

4 a 11 c 10 g 9 t

BASE COUNT
ORIGIN

Query Match 64.0%; Score 12.8; DB 17; Length 34;
Best Local Similarity 87.5%; Pred. No. 3.6e+04;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 GCGTGACGTGAGAG 20

11 | | | | | | | | | |

Db 32 GCGTGACGTGAGAG 17

RESULT 11
AA848120/c

LOCUS

DEFINITION
oe05a03.s1 NCI_CGAP_Ov2 Homo sapiens cDNA clone IMAGE:1384972
similar to gb:M35663 INTERFERON-INDUCED, DOUBLE-STRANDED
RNA-ACTIVATED PROTEIN KINASE (HUMAN);, mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

40 bp mRNA linear EST 07-APR-1998
AA848120
oe05a03.s1 NCI_CGAP_Ov2 Homo sapiens cDNA clone IMAGE:1384972
similar to gb:M35663 INTERFERON-INDUCED, DOUBLE-STRANDED
RNA-ACTIVATED PROTEIN KINASE (HUMAN);, mRNA sequence.
AA848120
AA848120.1 GI:2934638
human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 40)
NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapsb@mail.nih.gov
Tissue Procurement: Christopher A. Moskaluk, M.D., Michael R.
Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: David B. Krizman, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html

FEATURES
source

Trace considered overall poor quality
Insert Length: 1464 Std Error: 0.00
Seq primer: 40m13 fwd. ET from Amersham
High quality sequence stop: 1.
Location/Qualifiers
1. .40
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="IMAGE:1384972"
/clone_lib="NCI_CGAP_Ov2"
/sex="female"
/tissue_type="ovary"
/lab_host="DH10B"
/note="Vector: pAMP10; mRNA made from invasive ovarian
tumor, cDNA made by oligo-dT priming. Non-directionally
cloned. Size-selected on agarose gel, average insert size

600 bp. Reference: Krizman et al. (1996) Cancer Research
56:5380-5383."
8 a 14 c 9 g 9 t

Query Match 64.0%; Score 12.8; DB 9; Length 40;
Best Local Similarity 87.5%; Pred. No. 3.9e+04;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GATTGCGTCGCGTCAG 16

11 | | | | | | | | | |

Db 40 GATTGCGTCGCGTCAG 25

RESULT 12

AUI05994/c

LOCUS

DEFINITION
AUI05994 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone

KAIA2602, mRNA sequence.

ACCESSION
AUI05994

VERSION
AUI05994.1 GI:13555515

KEYWORDS
EST.

SOURCE
human.

ORGANISM
Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE
1 (bases 1 to 50)

AUTHORS
Suzuki,Y., Taira,H., Tsunoda,T., Mizushima-Sugano,J., Sese,J., Hata

,H., Ota,T., Isogai,T., Tanaka,T., Morishita,S., Okubo,K., Sakaki

,Y., Nakamura,Y., Suyama,A. and Sugano,S.

Diverse transcriptional initiation revealed by fine, large-scale

mapping of mRNA start sites

EMBO Rep. 2 (5), 388-393 (2001)

21270072

Contact: Yutaka Suzuki

Department of Virology

Institute of Medical Science, University of Tokyo

4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan

Email: ysuzuki@ims.u-tokyo.ac.jp

Suzuki,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and Sugano

,S. Construction and characterization of a full length-enriched and

a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).

Location/Qualifiers
1. .50
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="KAIA2602"
/note="Differential display comparison of untreated and
dimethylfumarate treated U937 cells"

BASE COUNT
11 a 19 c 9 g 11 t

ORIGIN

Query Match 64.0%; Score 12.8; DB 9; Length 50;
Best Local Similarity 87.5%; Pred. No. 4.3e+04;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GATTGCGTCGCGTCAG 16

11 | | | | | | | | | |

Db 29 GATTGCGTCGCGTCAG 14

RESULT 13

BH256492

LOCUS

DEFINITION
BH256492

KG03686-5prime Drosophila melanogaster P150P0r-P1 p element

insertion lines Drosophila melanogaster genomic Sequence recovered

from 5' end of P element, DNA sequence.

ACCESSION
BH256492

VERSION
BH256492.1 GI:17151385

KEYWORDS
GSS.

SOURCE
fruit fly.

ORGANISM
Drosophila melanogaster

Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

REFERENCE
AUTHORS
1 (bases 1 to 69)
G., Belen, H., Rubin, G. and Spradling, A.,
Tsang, G., He, Y., Karpen
Lewis, R., Hoskins, R., Liao, G., Mozen, N.,
The Berkeley Drosophila Genome Project
Unpublished (2001)

TITLE
JOURNAL
COMMENT
Contact: Gerald Rubin
Berkeley Drosophila Genome Project
University of California, Berkeley
LSA Building, Berkeley, CA 94720-3200, USA
Fax: 5106439947

Email: gerry@fruitfly.berkeley.edu
Sequence recovery method was inverse PCR.
Sequence orientation is forward strand relative to 5' end of P
element

The P element insertion position is base 62 in the 69 bases. This
insertion position refers to the first base of the 8 base target
recognition sequence.

Class: transposon-tagged.
Location/Qualifiers
1..69

FEATURES
source
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/clone_lib="Drosophila melanogaster P{SUPor-P} P element
insertion lines"

/note="Inverse PCR was performed on Drosophila
melanogaster strains each of which contains one or more
P{SUPor-P} P-element transposon insertion. The resultant
fragment for each strain was directly sequenced to
determine the genomic sequence at the site of insertion.
Details of the protocols used can be found at
<http://www.fruitfly.org/about/methods/inverse.pcr.html>."

BASE COUNT 15 a 15 c 18 g 21 t

ORIGIN

Query Match 64.0%; Score 12.8; DB 17; Length 69;
Best Local Similarity 87.5%; Pred. No. 4.9e+04;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ATTGCCTGAGCTCAGA 17
||||| ||| |||||
Db 2 ATTGCGTGAAGTCAGA 17

RESULT 14
AI204909
LOCUS
AI204909 86 bp mRNA linear EST 15-OCT-1998
an05g11.x1 Stratagene schizo brain S11 Homo sapiens cDNA clone
IMAGE:1684772 3' similar to gb:M55053 CYTOCHROME P450 1A2 (HUMAN);,
mRNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
AI204909.1 GI:3757971
EST.
Homo sapiens

ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE
AUTHORS
1 (bases 1 to 86)
Hillier, L., Allen, M., Bowles, L., Dubuque, T., Geisel, G., Jost, S.,
Krizman, D., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin
J., Moore, B., Schellenberg, K., Steptoe, M., Tan, F., Theising, B.,
White, Y., Wylie, T., Waterston, R. and Wilson, R.

TITLE
JOURNAL
COMMENT
Washington-NCI human EST Project
Unpublished (1997)
Contact: Wilson RK

Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810

Email: est@watson.wustl.edu
This clone is available royalty-free through LLNL ; contact the

IMAGE Consortium (info@image.llnl.gov) for further information.
Trace considered overall poor quality
Seq primer: -40UP from Gibco
High quality sequence stop: 1.

FEATURES
source
Location/Qualifiers
1..86
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="IMAGE:1684772"
/clone_lib="Stratagene schizo brain S11"
/sex="male"
/tissue_type="schizophrenic brain S-11 frontal lobe"
/dev_stage="34 years old"
/lab_host="SOLR (kanamycin resistant)"
/note="Vector: Bluescript SK-; Site:1: EcoRI; Library
constructed from S-11 frontal lobe, male, 34 years old,
50% caucasian, 50% Aleutian. Schizophrenic suicide.
Random primed into EcoRI site of ZAP II Vector. Mass
excised. Custom library. Avg insert length 1.4kb.
Material obtained by Johnston N., Torrey, E.F., Yolken R.,
and the Stanley Neuropathology Consortium - Analysis of
RNAs from the Brains of Individuals with Psychiatric
Diseases (Unpublished) Stanley Neurovirology Laboratory,
Johns Hopkins School of Medicine, Baltimore MD."

BASE COUNT 26 a 19 c 24 g 17 t

ORIGIN

Query Match 64.0%; Score 12.8; DB 9; Length 86;
Best Local Similarity 87.5%; Pred. No. 5.4e+04;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 GCCTGACGTCAGAGAG 20
||||| ||| |||||
Db 43 GCCTGCGCAGACAGAG 58

RESULT 15

AU260302/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

house mouse.

Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

Kato, K. and Matoba, R.

Generation of expressed

Unpublished (2002)

Contact: Kikuya Kato

Graduate School of Biological Sciences

Nara Institute of Science and Technology

8916-5 Takayama, Ikoma, Nara 630-0101, Japan

Tel: 81-743-72-5581

Fax: 81-743-72-5589

Email: kkatobs@ist-nara.ac.jp,

URL: <http://love2.aist-nara.ac.jp/BED/index.html>.

Location/Qualifiers

1..88

FEATURES

source

/organism="Mus musculus"

/db_xref="taxon:10090"

/clone="BED0016644"

/clone_lib="3'-directed mouse cDNA library"

/tissue_type="brain"

/note="Vector: pGEM-T-easy"

BASE COUNT 27 a 26 c 19 g 16 t

ORIGIN

Query Match

Best Local Similarity

Score 12.8; DB 9; Length 88;

Pred. No. 5.4e+04;

Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 GCGTGACGTCAGAGAG 20

|||||

Db 19 GCGTGAAGTCAGAGAG 4

Search completed: December 12, 2002, 06:03:49
Job time : 834.154 secs

GenCore version 5.1.3
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:12:31 ; Search time 33.8406 seconds
(without alignments)
233.720 Million cell updates/sec

Title: US-09-355-254F-8

Perfect score: 20
Sequence: 1 gattgcctgacgtcagag 20

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 355320 seqs, 197730502 residues

Total number of hits satisfying chosen parameters: 212722

Minimum DB seq length: 0
Maximum DB seq length: 100

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Published Applications_NA.*

- 1: /cgn2_6/ptodata/2/pubpna/US07_PUBCOMB.seq.*
- 2: /cgn2_6/ptodata/2/pubpna/US07_PUBCOMB.seq.*
- 3: /cgn2_6/ptodata/2/pubpna/US06_NEW_PUB.seq.*
- 4: /cgn2_6/ptodata/2/pubpna/US06_PUBCOMB.seq.*
- 5: /cgn2_6/ptodata/2/pubpna/US07_NEW_PUB.seq.*
- 6: /cgn2_6/ptodata/2/pubpna/US07_PUBCOMB.seq.*
- 7: /cgn2_6/ptodata/2/pubpna/US08_NEW_PUB.seq.*
- 8: /cgn2_6/ptodata/2/pubpna/US08_PUBCOMB.seq.*
- 9: /cgn2_6/ptodata/2/pubpna/US09_NEW_PUB.seq.*
- 10: /cgn2_6/ptodata/2/pubpna/US09_PUBCOMB.seq.*
- 11: /cgn2_6/ptodata/2/pubpna/US10_NEW_PUB.seq.*
- 12: /cgn2_6/ptodata/2/pubpna/US10_PUBCOMB.seq.*
- 13: /cgn2_6/ptodata/2/pubpna/US60_NEW_PUB.seq.*
- 14: /cgn2_6/ptodata/2/pubpna/US60_PUBCOMB.seq.*

pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20	100.0	27	10	US-09-878-918-2
2	19	95.0	23	10	US-09-816-763-148
3	12.8	64.0	50	10	US-09-875-945-4
4	12.8	64.0	97	10	US-09-864-761-19704
5	12.6	63.0	57	10	US-09-969-192-60
6	12.6	63.0	57	10	US-09-969-192-61
7	12.6	63.0	63	10	US-09-969-192-58
8	12.6	63.0	86	10	US-09-983-965-5019
9	12.2	61.0	18	10	US-09-420-433-62
10	12.2	61.0	24	10	US-09-784-911-30
11	12.2	61.0	99	10	US-09-769-066-11
12	12	60.0	63	10	US-09-998-598-2339
13	12	60.0	86	9	US-10-040-497-79
14	12	60.0	92	10	US-09-864-761-28968
15	11.8	59.0	18	8	US-08-983-605-140
16	11.6	58.0	18	10	US-09-303-510-50
17	11.6	58.0	18	10	US-09-303-040-50
18	11.6	58.0	20	9	US-10-060-301-123
19	11.6	58.0	41	10	US-09-238-351-29

c	20	11.6	58.0	41	10	US-09-238-351-40	Sequence 40, Appl
c	21	11.6	58.0	59	9	US-09-933-797-527	Sequence 527, App
	22	11.6	58.0	64	10	US-09-983-965-5569	Sequence 5569, Ap
	23	11.6	58.0	76	10	US-09-864-761-32057	Sequence 32057, A
	24	11.6	58.0	90	10	US-09-864-761-19844	Sequence 19844, A
	25	11.6	58.0	91	10	US-09-864-761-32056	Sequence 32056, A
	26	11.6	58.0	93	10	US-09-864-761-28341	Sequence 28341, A
c	27	11.6	58.0	98	10	US-09-864-761-26316	Sequence 26316, A
	28	11.4	57.0	23	10	US-09-893-238-22	Sequence 22, Appl
	29	11.4	57.0	47	9	US-09-853-526-242	Sequence 242, App
	30	11.4	57.0	47	10	US-09-901-484A-242	Sequence 242, App
	31	11.4	57.0	99	10	US-09-864-761-20908	Sequence 20908, A
	32	11.2	56.0	25	9	US-09-764-868-1251	Sequence 1251, Ap
	33	11.2	56.0	26	9	US-10-113-877-145	Sequence 145, App
c	34	11.2	56.0	31	10	US-09-801-274-276	Sequence 276, App
	35	11.2	56.0	44	10	US-09-875-519A-14	Sequence 14, Appl
c	36	11.2	56.0	47	9	US-09-853-526-319	Sequence 319, App
c	37	11.2	56.0	47	10	US-09-901-484A-319	Sequence 319, App
	38	11.2	56.0	59	10	US-09-875-945-12	Sequence 12, Appl
	39	11.2	56.0	84	10	US-09-764-860-1052	Sequence 1052, Ap
	40	11.2	56.0	84	10	US-09-764-860-1053	Sequence 1053, Ap
	41	11.2	56.0	84	10	US-09-764-877-3676	Sequence 3676, Ap
	42	11.2	56.0	88	10	US-09-764-869-1992	Sequence 1992, Ap
	43	11.2	56.0	88	10	US-09-764-869-1993	Sequence 1993, Ap
c	44	11.2	56.0	89	10	US-09-764-887-567	Sequence 567, App
	45	11.2	56.0	91	10	US-09-764-847-1964	Sequence 1964, Ap

ALIGNMENTS

RESULT 1
US-09-878-918-2
; Sequence 2, Application US/09878918
; Patent No. US00020107193A1
; GENERAL INFORMATION:
; APPLICANT: Glaxier Dr., Gordon W
; TITLE OF INVENTION: Therapeutic uses for IP3 receptor-mediated calcium channel modulators
; FILE REFERENCE: 84894-602
; CURRENT APPLICATION NUMBER: US/09/878,918
; CURRENT FILING DATE: 2002-04-03
; PRIOR APPLICATION NUMBER: 60/246763
; PRIOR FILING DATE: 2000-11-09
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2
; LENGTH: 27
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: CREB binding site
US-09-878-918-2

Query Match 100.0%; Score 20; DB 10; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.19;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTACAGAG 20
|||||
DB 4 GATTGCTGACGTACAGAG 23

RESULT 2
US-09-816-763-148
; Sequence 148, Application US/09816763
; Patent No. US20020110814A1
; GENERAL INFORMATION:
; APPLICANT: Remacle, Jose
; APPLICANT: Renard, Patricia
; APPLICANT: Art, Muriel
; TITLE OF INVENTION: METHOD AND KIT FOR THE SCREENING, THE

;; TITLE OF INVENTION: DETECTION AND/OR THE QUANTIFICATION OF TRANSCRIPTIONAL

;; FILE REFERENCE: VANM212.001AUS
;; CURRENT APPLICATION NUMBER: US/09/816,763
;; CURRENT FILING DATE: 2001-03-23
;; PRIOR APPLICATION NUMBER: EP 00870057.7
;; PRIOR FILING DATE: 2000-03-24
;; NUMBER OF SEQ ID NOS: 150
;; SOFTWARE: FastSeq for Windows version 4.0
;; SEQ ID NO 148
;; LENGTH: 23
;; TYPE: DNA
;; ORGANISM: Artificial Sequence
;; FEATURE:
;; OTHER INFORMATION: CREB consensus sequence
US-09-816-763-148

Query Match 95.0%; Score 19; DB 10; Length 23;
Best Local Similarity 100.0%; Pred. No. 0.62;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 ATTGCCTGACGTCAGAG 20

|||||

Db 1 ATTGCCTGACGTCAGAG 19

RESULT 3

US-09-875-945-4
;; Sequence 4, Application US/09875945
;; Patent No. US20020098169A1
;; GENERAL INFORMATION:
;; APPLICANT: METCON MEDICIN AB
;; APPLICANT: SMITH, ulf
;; TITLE OF INVENTION: No. US20020098169A1el sequences and their use
;; FILE REFERENCE: 45513MH
;; CURRENT APPLICATION NUMBER: US/09/875,945
;; CURRENT FILING DATE: 2001-06-08
;; PRIOR APPLICATION NUMBER: SE 0002189-9
;; PRIOR FILING DATE: 2000-06-09
;; PRIOR APPLICATION NUMBER: US 60/210,207
;; PRIOR FILING DATE: 2000-06-08
;; NUMBER OF SEQ ID NOS: 14
;; SOFTWARE: PatentIn Ver. 2.1
;; SEQ ID NO 4
;; TYPE: DNA
;; LENGTH: 50
;; ORGANISM: Homo sapiens
US-09-875-945-4

Query Match 64.0%; Score 12.8; DB 10; Length 50;

Best Local Similarity 87.5%; Pred. No. 9.4e+02;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAG 16

|||||

Db 6 GATTGCTGACGTCAG 21

RESULT 4

US-09-864-761-19704
;; Sequence 19704, Application US/09864761
;; Patent No. US20020048763A1
;; GENERAL INFORMATION:
;; APPLICANT: Penn, Sharron G.
;; APPLICANT: Rank, David R.
;; APPLICANT: Hanzel, David K.
;; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
;; TITLE OF INVENTION: GENE EXPRESSION ANALYSIS BY MICROARRAY
;; FILE REFERENCE: Aeonica-X-1
;; CURRENT APPLICATION NUMBER: US/09/864,761
;; CURRENT FILING DATE: 2001-05-23
;; PRIOR APPLICATION NUMBER: US 60/180,312

;; PRIOR FILING DATE: 2000-02-04
;; PRIOR APPLICATION NUMBER: US 60/207,456
;; PRIOR FILING DATE: 2000-05-26
;; PRIOR APPLICATION NUMBER: US 09/632,366
;; PRIOR FILING DATE: 2000-08-03
;; PRIOR APPLICATION NUMBER: GB 24263.6
;; PRIOR FILING DATE: 2000-10-04
;; PRIOR APPLICATION NUMBER: US 60/236,359
;; PRIOR FILING DATE: 2000-09-27
;; PRIOR APPLICATION NUMBER: PCT/US01/006666
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00667
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00664
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00669
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00665
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00668
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00663
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00662
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00661
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00670
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: US 60/234,687
;; PRIOR FILING DATE: 2000-09-21
;; PRIOR APPLICATION NUMBER: US 09/608,408
;; PRIOR FILING DATE: 2000-06-30
;; PRIOR APPLICATION NUMBER: US 09/774,203
;; PRIOR FILING DATE: 2001-01-29
;; NUMBER OF SEQ ID NOS: 49117
;; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
;; SEQ ID NO 19704
;; LENGTH: 97
;; TYPE: DNA
;; ORGANISM: Homo sapiens
;; FEATURE:
;; OTHER INFORMATION: MAP TO AC009490.4
;; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.6
;; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 1.8
;; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.5
;; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 1.7
;; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 1.5
;; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1.9
;; OTHER INFORMATION: EXPRESSED IN BT474, SIGNAL = 1.6
;; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 2
;; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 1.7
;; OTHER INFORMATION: EXPRESSED IN HBL100, SIGNAL = 1.7
;; OTHER INFORMATION: NT HIT: X56160.1, EVALUAE 3.20e+00
;; OTHER INFORMATION: EST_HUMAN HIT: AV648669.1, EVALUAE 1.90e-02
US-09-864-761-19704

Query Match 64.0%; Score 12.8; DB 10; Length 97;
Best Local Similarity 87.5%; Pred. No. 1e+03;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 GCCTGACGTCAGAG 20

|||||

Db 74 GCAGACATCAGAG 89

RESULT 5

US-09-969-192-60
;; Sequence 60, Application US/09969192
;; Patent No. US20020151027A1
;; GENERAL INFORMATION:
;; APPLICANT: WICKHAM, THOMAS J.
;; APPLICANT: ROELVINK, PETRUS W.

;; KOVESDI, IMRE
;; TITLE OF INVENTION: TARGETING ADENOVIRUS WITH USE OF
;; CONSTRAINED PEPTIDE MOTIFS
;;
;; NUMBER OF SEQUENCES: 80
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: Leydig, Voit & Mayer, Ltd.
;; STREET: Two Prudential Plaza - 49th Floor
;; CITY: Chicago
;; STATE: Illinois
;; COUNTRY: USA
;; ZIP: 60601
;;
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: PatentIn Release #1.0, Version #1.30
;;
;; CURRENT APPLICATION DATA: US/09/969,192
;; FILING DATE: 01-Oct-2001
;;
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 9-455061
;; FILING DATE: 06-DEC-1999
;; APPLICATION NUMBER: US 9-130225
;; FILING DATE: 06-AUG-1998
;; APPLICATION NUMBER: US 8-701124
;; FILING DATE: 21-AUG-1996
;;
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Hefner, M. Daniel
;; REGISTRATION NUMBER: 41,826
;; REFERENCE/DOCKET NUMBER: 213564
;;
;; INFORMATION FOR SEQ ID NO: 60:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 57 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;;
;; MOLECULE TYPE: other nucleic acid
;; SEQUENCE DESCRIPTION: SEQ ID NO: 60:
US-09-969-192-60

Query Match 63.0%; Score 12.6; DB 10; Length 57;
Best Local Similarity 78.9%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 ATTGCCTGACGTCAGAG 20
| | | | | | | | | |
Db 8 AATTCTTGACGTCGAG 26

RESULT 6
US-09-969-192-61/c
; Sequence 61, Application US/09969192
; Patent No. US20020151027A1
; GENERAL INFORMATION:
; APPLICANT: WICKHAM, THOMAS J.
; ROELVINK, PETRUS W.
; KOVESDI, IMRE
; TITLE OF INVENTION: TARGETING ADENOVIRUS WITH USE OF
; CONSTRAINED PEPTIDE MOTIFS
;
; NUMBER OF SEQUENCES: 80
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Leydig, Voit & Mayer, Ltd.
; STREET: Two Prudential Plaza - 49th Floor
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60601
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:

;; APPLICATION NUMBER: US/09/969,192
;; FILING DATE: 01-Oct-2001
;;
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 9-455061
;; FILING DATE: 06-DEC-1999
;; APPLICATION NUMBER: US 9-130225
;; FILING DATE: 06-AUG-1998
;; APPLICATION NUMBER: US 8-701124
;; FILING DATE: 21-AUG-1996
;;
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Hefner, M. Daniel
;; REGISTRATION NUMBER: 41,826
;; REFERENCE/DOCKET NUMBER: 213564
;;
;; INFORMATION FOR SEQ ID NO: 61:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 57 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;;
;; MOLECULE TYPE: other nucleic acid
;; SEQUENCE DESCRIPTION: SEQ ID NO: 61:
US-09-969-192-61

Query Match 63.0%; Score 12.6; DB 10; Length 57;
Best Local Similarity 78.9%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 ATTGCCTGACGTCAGAG 20
| | | | | | | | | |
Db 54 AATTCTTGACGTCGAG 36

RESULT 7
US-09-969-192-58
; Sequence 58, Application US/09969192
; Patent No. US20020151027A1
; GENERAL INFORMATION:
; APPLICANT: WICKHAM, THOMAS J.
; ROELVINK, PETRUS W.
; KOVESDI, IMRE
; TITLE OF INVENTION: TARGETING ADENOVIRUS WITH USE OF
; CONSTRAINED PEPTIDE MOTIFS
;
; NUMBER OF SEQUENCES: 80
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Leydig, Voit & Mayer, Ltd.
; STREET: Two Prudential Plaza - 49th Floor
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60601
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/969,192
; FILING DATE: 01-Oct-2001
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 9-455061
; FILING DATE: 06-DEC-1999
; APPLICATION NUMBER: US 9-130225
; FILING DATE: 06-AUG-1998
; APPLICATION NUMBER: US 8-701124
; FILING DATE: 21-AUG-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Hefner, M. Daniel
; REGISTRATION NUMBER: 41,826
; REFERENCE/DOCKET NUMBER: 213564
; INFORMATION FOR SEQ ID NO: 58:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 63 base pairs
; TYPE: nucleic acid

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;
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; SEQUENCE DESCRIPTION: SEQ ID NO: 58:
US-09-969-192-58
Query Match 63.0%; Score 12.6; DB 10; Length 63;
Best Local Similarity 78.9%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 ATTGCTGACGTCAGAG 20
Db 9 AATTCTTGACGTCGGAG 27

RESULT 8
US-09-983-965-5019/c
; Sequence 5019, Application US/09983965
; Patent No. US20020137160A1
; GENERAL INFORMATION:
; APPLICANT: Warren, Wesley C.
; APPLICANT: Tao, Nengbing
; APPLICANT: Byatt, John C.
; APPLICANT: Mathialagan, Nagappan
; TITLE OF INVENTION: NUCLEIC ACID AND OTHER MOLECULES ASSOCIATED WITH LACTATION AND
; FILE REFERENCE: 37-21(10297)C
; CURRENT APPLICATION NUMBER: US/09/983,965
; CURRENT FILING DATE: 2001-10-26
; PRIOR APPLICATION NUMBER: US 09/465,231
; PRIOR FILING DATE: 1999-12-15
; PRIOR APPLICATION NUMBER: US 60/113,678
; PRIOR FILING DATE: 1998-12-17
; NUMBER OF SEQ ID NOS: 5912
; SEQ ID NO 5019
; LENGTH: 86
; TYPE: DNA
; ORGANISM: Bos taurus
; FEATURE:
; OTHER INFORMATION: Clone ID: 29-LIB34-022-Q1-E1-H1
US-09-983-965-5019

Query Match 63.0%; Score 12.6; DB 10; Length 86;
Best Local Similarity 78.9%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAGA 19
Db 61 GATGGCTGGGGTCAAGA 43

RESULT 9
US-09-420-433-62
; Sequence 62, Application US/09420433
; Patent No. US20020096480A1
; GENERAL INFORMATION:
; APPLICANT: Sidransky, David
; TITLE OF INVENTION: NUCLEIC ACID MUTATION DETECTION IN
; TITLE OF INVENTION: HISTOLOGIC TISSUE
; NUMBER OF SEQUENCES: 82
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Spensley Horn Jubas & Lubitz
; STREET: 1880 Century Park East, Suite 500
; CITY: Los Angeles
; STATE: California
; COUNTRY: USA
; ZIP: 90067
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
```

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;
; APPLICATION NUMBER: US/09/420.433
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/181.664
; FILING DATE: JANUARY 14, 1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Wetherell, Jr., Ph.D., John R.
; REGISTRATION NUMBER: 31,678
; REFERENCE/DOCKET NUMBER: PD-3055
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 455-5100
; TELEFAX: (619) 455-5110
; INFORMATION FOR SEQ ID NO: 62:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 18 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..18
US-09-420-433-62

Query Match 61.0%; Score 12.2; DB 10; Length 18;
Best Local Similarity 82.4%; Pred. No. 1.7e+03;
Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 4 TGCCTGACGTCAGAG 20
Db 1 TGCCTGTCGGGAG 17

RESULT 10
US-09-784-911-30/c
; Sequence 30, Application US/09784911
; Patent No. US20020072115A1
; GENERAL INFORMATION:
; APPLICANT: Harrison, Leonard C.
; APPLICANT: Jiang, Fang-Xu
; APPLICANT: Stanley, Edouard Guy
; APPLICANT: Gopez, Leonel Jorge
; TITLE OF INVENTION: Pancreatic islet cell growth factors
; FILE REFERENCE: Davies Collison Cave
; CURRENT APPLICATION NUMBER: US/09/784,911
; CURRENT FILING DATE: 2001-09-17
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: PatentIn version 2.1
; SEQ ID NO 30
; LENGTH: 24
; TYPE: DNA
; ORGANISM: primer
US-09-784-911-30

Query Match 61.0%; Score 12.2; DB 10; Length 24;
Best Local Similarity 82.4%; Pred. No. 1.8e+03;
Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 4 TGCCTGACGTCAGAG 20
Db 17 TGCCTGACATCAAGAG 1

RESULT 11
US-09-769-066-11
; Sequence 11, Application US/09769066
; Patent No. US20020107360A1
; GENERAL INFORMATION:
; APPLICANT: Fuerst, Thomas R.
; McAttee, C. Patrick
; Yarbrough, Patrice O.
; Zhang, Yifan
```

;; TITLE OF INVENTION: HEPATITIS E VIRUS ANTIGENS AND USES THEREFOR
;; NUMBER OF SEQUENCES: 31
;; CORRESPONDENCE ADDRESS:

;; ADDRESSEE: Dehlinger & Associates
;; STREET: 350 Cambridge Ave., Suite 250
;; CITY: Palo Alto
;; STATE: CA
;; COUNTRY: USA

;; ZIP: 94306

;; COMPUTER READABLE FORM:

;; MEDIUM TYPE: Floppy disk

;; COMPUTER: IBM PC compatible

;; OPERATING SYSTEM: PC-DOS/MS-DOS

;; SOFTWARE: Patent In Release #1.0, Version #1.25

;; CURRENT APPLICATION DATA:

;; APPLICATION NUMBER: US/09/769,066

;; FILING DATE: 24-Jan-2001

;; CLASSIFICATION: <Unknown>

;; PRIOR APPLICATION DATA:

;; APPLICATION NUMBER: 08/542,634

;; FILING DATE: <Unknown>

;; ATTORNEY/AGENT INFORMATION:

;; NAME: Fabian, Gary R.

;; REGISTRATION NUMBER: 33,875

;; REFERENCE/DOCKET NUMBER: 4600-0293.30

;; TELECOMMUNICATION INFORMATION:

;; TELEPHONE: (415) 324-0880

;; TELEFAX: (415) 324-0960

;; INFORMATION FOR SEQ ID NO: 11:

;; SEQUENCE CHARACTERISTICS:

;; LENGTH: 99 base pairs

;; TYPE: nucleic acid

;; STRANDEDNESS: Hepatitis E Virus (Burma strain)

;; 406.4-2 region

;; TOPOLOGY: linear

;; MOLECULE TYPE: DNA (genomic)

;; HYPOTHETICAL: NO

;; ORIGINAL SOURCE:

;; SEQUENCE DESCRIPTION: SEQ ID NO: 11:

US-09-769-066-11

Query Match 61.0%; Score 12.2; DB 10; Length 99;

Best Local Similarity 82.4%; Pred. No. 2e+03;

Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 TTGCCTGACGTCAGAGA 19

||||| ||||| |||

DB 58 TTGCCTGACGTCGTAGA 74

RESULT 12

US-09-998-598-2339

;; Sequence 2339, Application US/09998598

;; Patent No. US20020150922A1

;; GENERAL INFORMATION:

;; APPLICANT: Stolk, John A.

;; APPLICANT: Xu, Jiangchun

;; APPLICANT: Chenault, Ruth A.

;; APPLICANT: Meagher, Madelein Joy

;; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND

;; FILE REFERENCE: 210121.561

;; CURRENT APPLICATION NUMBER: US/09/998,598

;; NUMBER OF SEQ ID NOS: 2606

;; SOFTWARE: Corixa Invention Disclosure Database

;; SEQ ID NO 2339

;; LENGTH: 63

;; TYPE: DNA

;; ORGANISM: Homo sapiens

US-09-998-598-2339

Query Match 60.0%; Score 12; DB 10; Length 63;

Best Local Similarity 75.0%; Pred. No. 2.4e+03;

Matches 15; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAG 20

|| ||||| ||||| |||

DB 14 GAACTCTGACGTCAAAGTG 33

RESULT 13

US-10-040-497-79

;; Sequence 79, Application US/10040497

;; Patent No. US20020172962A1

;; GENERAL INFORMATION:

;; APPLICANT: GOLD, LARRY

;; TUEK, CRAIG

;; TITLE OF INVENTION: METHODS OF PRODUCING NUCLEIC ACID LIGANDS

;; NUMBER OF SEQUENCES: 83

;; CORRESPONDENCE ADDRESS:

;; ADDRESSEE: Swanson & Bratschun, L.L.C.

;; STREET: 8400 E. Prentice Avenue, Suite 200

;; CITY: Englewood

;; STATE: Colorado

;; COUNTRY: USA

;; ZIP: 80111

;; COMPUTER READABLE FORM:

;; MEDIUM TYPE: Diskette, 3 1/5 inch, 1.44 MB

;; COMPUTER: IBM compatible

;; OPERATING SYSTEM: MS-DOS

;; SOFTWARE: WordPerfect 8.0

;; CURRENT APPLICATION DATA:

;; APPLICATION NUMBER: US/10/040,497

;; FILING DATE: 07-Jan-2002

;; PRIOR APPLICATION DATA:

;; APPLICATION NUMBER: 08/748,697

;; FILING DATE: 13-NOVEMBER-1996

;; APPLICATION NUMBER: 08/442,062

;; FILING DATE: 16-MAY-1995

;; APPLICATION NUMBER: 07/964,624

;; FILING DATE: 21-OCTOBER-1992

;; APPLICATION NUMBER: 07/714,131

;; FILING DATE: 10-JUNE-1991

;; APPLICATION NUMBER: 07/536,428

;; FILING DATE: 11-JUNE-1990

;; ATTORNEY/AGENT INFORMATION:

;; NAME: Barry J. Swanson

;; REGISTRATION NUMBER: 33,215

;; REFERENCE/DOCKET NUMBER: NEX05/DC-CON

;; TELECOMMUNICATION INFORMATION:

;; TELEPHONE: (303) 793-3333

;; TELEFAX: (303) 793-3433

;; INFORMATION FOR SEQ ID NO: 79:

;; SEQUENCE CHARACTERISTICS:

;; LENGTH: 86 base pairs

;; TYPE: nucleic acid

;; STRANDEDNESS: single

;; TOPOLOGY: linear

;; SEQUENCE DESCRIPTION: SEQ ID NO: 79:

US-10-040-497-79

Query Match

Best Local Similarity 60.0%; Score 12; DB 9; Length 86;

Matches 13; Conservative 2; Mismatches 5; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAG 20

|| ||||| || |||||

DB 10 GAUGGCCUCCGACGAG 29

RESULT 14

US-09-864-761-28968

;; Sequence 28968, Application US/09864761

;; Patent No. US20020048763A1

;; GENERAL INFORMATION:


```
; EARLIER FILING DATE: 1998-08-12
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 10
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Chlamydia trachomatis
; FEATURE:
; OTHER INFORMATION: An oligonucleotide derived from the DNA encoding a
; OTHER INFORMATION: 60 kDa cysteine rich outer membrane protein from
; OTHER INFORMATION: Chlamydia trachomatis.
US-09-303-862-10

Query Match          100.0%; Score 20; DB 3; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.057;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAG 20
Db 1 GATTGCTGACGTCAGAG 20

RESULT 3
US-08-210-880B-2
; Sequence 2, Application US/08210880B
; Patent No. 5641486
; GENERAL INFORMATION:
; APPLICANT: HINRICHS, STEVEN H.
; ATTORNEY/AGENT INFORMATION:
; TITLE OF INVENTION: METHOD AND COMPOSITION FOR INHIBITING
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HENDERSON & STURM
; STREET: 1125 S. 103RD ST., #330
; CITY: OMAHA
; STATE: NE
; COUNTRY: US
; ZIP: 68124
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; FILING DATE: 18-MAR-1994
; APPLICATION NUMBER: US/08/210,880B
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: JONDLE, ROBERT J.
; REGISTRATION NUMBER: 33,915
; REFERENCE/DOCKET NUMBER: 63066
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 402-398-9000
; TELEFAX: 402-398-9005
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 27 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: unknown
; MOLECULE TYPE: DNA (genomic)
US-08-210-880B-2

Query Match          100.0%; Score 20; DB 1; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.059;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAG 20
Db 4 GATTGCTGACGTCAGAG 23

US-08-210-880B-2

; EARLIER FILING DATE: 1998-08-12
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 10
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Chlamydia trachomatis
; FEATURE:
; OTHER INFORMATION: An oligonucleotide derived from the DNA encoding a
; OTHER INFORMATION: 60 kDa cysteine rich outer membrane protein from
; OTHER INFORMATION: Chlamydia trachomatis.
US-09-303-862-10

Query Match          100.0%; Score 20; DB 3; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.057;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAG 20
Db 1 GATTGCTGACGTCAGAG 20

RESULT 4
US-08-771-411-2
; Sequence 2, Application US/08771411
; Patent No. 5844096
; GENERAL INFORMATION:
; APPLICANT: HINRICHS, STEVEN H.
; ATTORNEY/AGENT INFORMATION:
; TITLE OF INVENTION: METHOD AND COMPOSITION FOR INHIBITING
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HENDERSON & STURM
; STREET: 1125 S. 103RD ST., #330
; CITY: OMAHA
; STATE: NE
; COUNTRY: US
; ZIP: 68124
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; FILING DATE: 20-DEC-1996
; APPLICATION NUMBER: US/08/771,411
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; FILING DATE: 18-MAR-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: JONDLE, ROBERT J.
; REGISTRATION NUMBER: 33,915
; REFERENCE/DOCKET NUMBER: 63066
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 402-398-9000
; TELEFAX: 402-398-9005
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 27 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: unknown
; MOLECULE TYPE: DNA (genomic)
US-08-771-411-2

Query Match          100.0%; Score 20; DB 2; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.059;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAG 20
Db 4 GATTGCTGACGTCAGAG 23

RESULT 5
US-09-215-098-1
; Sequence 1, Application US/09215098
; Patent No. 6194632
; GENERAL INFORMATION:
; APPLICANT: Leiden, Jeffery M
; TITLE OF INVENTION: DILATED CARDIOMYOPATHY IN TRANSGENIC MICE EXPRESSING A
; TITLE OF INVENTION: DOMINANT-NEGATIVE CREB TRANSCRIPTION FACTOR IN THE
; TITLE OF INVENTION: HEART
; FILE REFERENCE: 9189-4
; CURRENT APPLICATION NUMBER: US/09/215,098
; CURRENT FILING DATE: 1998-12-18
; PRIOR APPLICATION NUMBER: 60/068,011
; PRIOR FILING DATE: 1997-12-18
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 32
; TYPE: DNA
```

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; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence:Oligonucleotide
; OTHER INFORMATION: containing the CREB site from the somatostatin
; OTHER INFORMATION: promoter
US-09-215-098-1

Query Match      82.0%; Score 16.4; DB 4; Length 32;
Best Local Similarity 94.4%; Pred. No. 4.9;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 TTGCCTGACGTCAGAG 20
   ||| ||||| ||||| |||||
Db 11 TTGCCTGACGTCAGAG 28

RESULT 6
US-08-171-389-451
; Sequence 451, Application US/08171389
; Patent No. 5578444
; GENERAL INFORMATION:
; APPLICANT: Edwards, Cynthia A.
; APPLICANT: Cantor, Charles R.
; APPLICANT: Andrews, Beth M.
; APPLICANT: Turin, Lisa M.
; APPLICANT: Fry, Kirk E.
; TITLE OF INVENTION: Sequence-Directed DNA Binding
; TITLE OF INVENTION: Molecules, Compositions and Methods
; NUMBER OF SEQUENCES: 641
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genelabs Technologies, Inc.
; STREET: 505 Penobscot Drive
; CITY: Redwood City
; STATE: CA
; COUNTRY: USA
; ZIP: 94063
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/171,389
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/123,936
; FILING DATE: 17-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/996,783
; FILING DATE: 23-DEC-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/723,618
; FILING DATE: 27-JUN-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/081,070
; FILING DATE: 22-JUN-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Fabian, Gary R.
; REGISTRATION NUMBER: 33,875
; REFERENCE/DOCKET NUMBER: 4600-0175/G19P3
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 324-0880
; TELEFAX: (415) 324-0960
; INFORMATION FOR SEQ ID NO: 451:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 50 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ORIGINAL SOURCE:
; INDIVIDUAL ISOLATE: Human somatostatin I gene
US-08-123-936-451

Query Match      82.0%; Score 16.4; DB 1; Length 50;
Best Local Similarity 94.4%; Pred. No. 5.2;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
; INDIVIDUAL ISOLATE: Human somatostatin I gene
US-08-171-389-451

Query Match      82.0%; Score 16.4; DB 1; Length 50;
Best Local Similarity 94.4%; Pred. No. 5.2;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 TTGCCTGACGTCAGAG 20
   ||| ||||| ||||| |||||
Db 1 TAGCCTGACGTCAGAG 18

RESULT 7
US-08-123-936-451
; Sequence 451, Application US/08123936
; Patent No. 5726014
; GENERAL INFORMATION:
; APPLICANT: Edwards, Cynthia A.
; APPLICANT: Cantor, Charles R.
; APPLICANT: Andrews, Beth M.
; APPLICANT: Turin, Lisa M.
; TITLE OF INVENTION: Screening Assay for the Detection of
; TITLE OF INVENTION: DNA-Binding Molecules
; NUMBER OF SEQUENCES: 640
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genelabs Technologies, Inc.
; STREET: 505 Penobscot Drive
; CITY: Redwood City
; STATE: CA
; COUNTRY: USA
; ZIP: 94063
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/123,936
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/996,783
; FILING DATE: 23-DEC-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/723,618
; FILING DATE: 27-JUN-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Fabian, Gary R.
; REGISTRATION NUMBER: 33,875
; REFERENCE/DOCKET NUMBER: 4600-0075.32/G19P2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 324-0880
; TELEFAX: (415) 324-0960
; INFORMATION FOR SEQ ID NO: 451:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 50 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ORIGINAL SOURCE:
; INDIVIDUAL ISOLATE: Human somatostatin I gene
US-08-123-936-451

Query Match      82.0%; Score 16.4; DB 1; Length 50;
Best Local Similarity 94.4%; Pred. No. 5.2;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 TTGCCTGACGTCAGAG 20
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Db 1 TAGCCTGACGTCAGAG 18
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RESULT 8
US-08-475-228A-451
; Sequence 451, Application US/08475228A
; Patent No. 5869241
; GENERAL INFORMATION:
; APPLICANT: Edwards, Cynthia A.
; APPLICANT: Cantor, Charles R.
; APPLICANT: Andrews, Beth M.
; APPLICANT: Turin, Lisa M.
; APPLICANT: Fry, Kirk E.
; TITLE OF INVENTION: Sequence-Directed DNA Binding
; TITLE OF INVENTION: Molecules, Compositions and Methods
; NUMBER OF SEQUENCES: 664
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genelabs Technologies, Inc.
; STREET: 505 Penobscot Drive
; CITY: Redwood City
; STATE: CA
; COUNTRY: USA
; ZIP: 94063
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/475,228A
; FILING DATE: 06-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/123,936
; FILING DATE: 17-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/996,783
; FILING DATE: 23-DEC-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/723,618
; FILING DATE: 27-JUN-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/081,070
; FILING DATE: 22-JUN-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Stratford, Carol A.
; REGISTRATION NUMBER: 34,444
; REFERENCE/DOCKET NUMBER: 4600-0175.21/G19P3D2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 324-0880
; TELEFAX: (415) 324-0960
; INFORMATION FOR SEQ ID NO: 451:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 50 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ORIGINAL SOURCE:
; INDIVIDUAL ISOLATE: Human somatostatin I gene
US-08-475-228A-451

Query Match 82.0%; Score 16.4; DB 2; Length 50;
Best Local Similarity 94.4%; Pred. NO. 5.2;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 TTGCCTGACGTCAGAG 20
Db 1 TAGCCTGACGTCAGAG 18

RESULT 9
US-08-482-080A-451
; Sequence 451, Application US/08482080A
; Patent No. 6010849

; GENERAL INFORMATION:
; APPLICANT: Edwards, Cynthia A.
; APPLICANT: Cantor, Charles R.
; APPLICANT: Andrews, Beth M.
; APPLICANT: Turin, Lisa M.
; APPLICANT: Fry, Kirk E.
; TITLE OF INVENTION: Sequence-Directed DNA Binding
; TITLE OF INVENTION: Molecules, Compositions and Methods
; NUMBER OF SEQUENCES: 664
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genelabs Technologies, Inc.
; STREET: 505 Penobscot Drive
; CITY: Redwood City
; STATE: CA
; COUNTRY: USA
; ZIP: 94063
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/482,080A
; FILING DATE: 07-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/171,389
; FILING DATE: 20-DEC-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/123,936
; FILING DATE: 17-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/996,783
; FILING DATE: 23-DEC-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/723,618
; FILING DATE: 27-JUN-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/081,070
; FILING DATE: 22-JUN-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Brady, John F.
; REGISTRATION NUMBER: 39,118
; REFERENCE/DOCKET NUMBER: 4600-0175.20/G19P3D1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650) 324-0880
; TELEFAX: (650) 324-0960
; INFORMATION FOR SEQ ID NO: 451:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 50 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ORIGINAL SOURCE:
; INDIVIDUAL ISOLATE: Human somatostatin I gene
US-08-482-080A-451

Query Match 82.0%; Score 16.4; DB 3; Length 50;
Best Local Similarity 94.4%; Pred. NO. 5.2;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 TTGCCTGACGTCAGAG 20
Db 1 TAGCCTGACGTCAGAG 18

RESULT 10
US-09-354-947-451
; Sequence 451, Application US/09354947
; Patent No. 6384208
; GENERAL INFORMATION:
; APPLICANT: Edwards, Cynthia A.

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; APPLICANT: Cantor, Charles R.
; APPLICANT: Andrews, Beth M.
; APPLICANT: Turin, Lisa M.
; APPLICANT: Fry, Kirk E.
; TITLE OF INVENTION: Sequence-Directed DNA Binding
; NUMBER OF SEQUENCES: 664
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genelabs Technologies, Inc.
; STREET: 505 Penobscot Drive
; CITY: Redwood City
; STATE: CA
; COUNTRY: USA
; ZIP: 94063
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/354,947
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/482,080
; FILING DATE: 07-JUN-1995
; APPLICATION NUMBER: US 08/171,389
; FILING DATE: 20-DEC-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/123,936
; FILING DATE: 17-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/996,783
; FILING DATE: 23-DEC-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/081,070
; FILING DATE: 22-JUN-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Brady, John F.
; REGISTRATION NUMBER: 39,118
; REFERENCE/DOCKET NUMBER: 4600-0175.20/G19P3D1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650) 324-0880
; TELEFAX: (650) 324-0960
; INFORMATION FOR SEQ ID NO: 451:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 50 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ORIGINAL SOURCE:
; INDIVIDUAL ISOLATE: Human somatostatin I gene
; US-09-354-947-451

Query Match      82.0%; Score 16.4; DB 4; Length 50;
Best Local Similarity 94.4%; Pred. No. 5.2;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY  3 TTGCCTGACGTCAGAG 20
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Db   1 TAGCCTGACGTCAGAG 18

RESULT 11
PCT-US93-12388-451
; Sequence 451, Application PC/TUS9312388
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: Sequence-Directed DNA Binding

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; TITLE OF INVENTION: Molecules, Compositions and Methods
; NUMBER OF SEQUENCES: 641
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genelabs Technologies, Inc.
; STREET: 505 Penobscot Drive
; CITY: Redwood City
; STATE: CA
; COUNTRY: USA
; ZIP: 94063
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US93/12388
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/123,936
; FILING DATE: 17-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/996,783
; FILING DATE: 23-DEC-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Fabian, Gary R.
; REGISTRATION NUMBER: 33,875
; REFERENCE/DOCKET NUMBER: 4600-0175.41/G19PCT2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 324-0880
; TELEFAX: (415) 324-0960
; INFORMATION FOR SEQ ID NO: 451:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 50 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ORIGINAL SOURCE:
; INDIVIDUAL ISOLATE: Human somatostatin I gene
; PCT-US93-12388-451

Query Match      82.0%; Score 16.4; DB 5; Length 50;
Best Local Similarity 94.4%; Pred. No. 5.2;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY  3 TTGCCTGACGTCAGAG 20
    | | | | | | | | | |
Db   1 TAGCCTGACGTCAGAG 18

RESULT 12
US-08-813-507-78/c
; Sequence 78, Application US/08813507
; Patent No. 6114116
; GENERAL INFORMATION:
; APPLICANT: Lemieux, Bertrand
; APPLICANT: Landry, Benoit S.
; APPLICANT: Sapolsky, Ronald J.
; TITLE OF INVENTION: Brassica Polymorphisms
; NUMBER OF SEQUENCES: 173
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, Eighth Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS

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SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/813,507
FILING DATE: 07-MAR-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/032,069
FILING DATE: 02-DEC-1996
ATTORNEY/AGENT INFORMATION:
NAME: Liebeschuetz, Joe
REGISTRATION NUMBER: 37,505
REFERENCE/DOCKET NUMBER: 018547-030100US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415 576-0200
TELEFAX: 415 576-0200
TELEX:
INFORMATION FOR SEQ ID NO: 78:
SEQUENCE CHARACTERISTICS:
LENGTH: 41 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-813-507-78

Query Match 71.0%; Score 14.2; DB 3; Length 41;
Best Local Similarity 84.2%; Pred. No. 74;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAGA 19
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Db 19 GCTTTCATGACGTCAGAGA 1

RESULT 13

US-09-464-453-78/c
Sequence 78, Application US/09464453
Patent No. 6358686
GENERAL INFORMATION:
APPLICANT: Lemieux, Bertrand
Sepolsky, Ronald J.
TITLE OF INVENTION: Brassinica Polymorphisms
NUMBER OF SEQUENCES: 173
CORRESPONDENCE ADDRESS:
ADDRESS: Townsend and Townsend and Crew LLP
STREET: Two Embarcadero Center, Eighth Floor
CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94111-3834
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/464,453
FILING DATE: 14-DEC-1999
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/813,507
FILING DATE: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Liebeschuetz, Joe
REGISTRATION NUMBER: 37,505
REFERENCE/DOCKET NUMBER: 018547-030100US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415 576-0200
TELEFAX: 415 576-0200
TELEX: <Unknown>
INFORMATION FOR SEQ ID NO: 78:
SEQUENCE CHARACTERISTICS:
LENGTH: 41 base pairs

TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 78:
US-09-464-453-78

Query Match 71.0%; Score 14.2; DB 4; Length 41;
Best Local Similarity 84.2%; Pred. No. 74;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAGA 19
| | | | | | | | | | | | | | | | | | | | | |
Db 19 GCTTTCATGACGTCAGAGA 1

RESULT 14

US-08-870-930-25
Sequence 25, Application US/08870930
Patent No. 6168778
GENERAL INFORMATION:
APPLICANT: NEBOJSA JANJIC, LARRY GOLD, PAUL G. SCHMIDT, CHANDRA VARGESE, MICHA
TITLE OF INVENTION: VASCULAR ENDOTHELIAL GROWTH FACTOR (VEGF)
NUMBER OF SEQUENCES: 88
CORRESPONDENCE ADDRESS:
ADDRESS: Swanson and Bratschun, L.L.C.
STREET: 8400 East Prentice Avenue, Suite #200
CITY: Denver
STATE: Colorado
COUNTRY: USA
ZIP: 80111

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 1.4 Mb storage
COMPUTER: IBM compatible
OPERATING SYSTEM: MS-DOS
SOFTWARE: WordPerfect 8.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/870,930
FILING DATE: 6 JUNE 1997
CLASSIFICATION: 536

ATTORNEY/AGENT INFORMATION:
NAME: Barry J. Swanson
REGISTRATION NUMBER: 33,215
REFERENCE/DOCKET NUMBER: NEX61
TELECOMMUNICATION INFORMATION:
TELEPHONE: (303) 793-3333
TELEFAX: (303) 793-3433
INFORMATION FOR SEQ ID NO: 25:
SEQUENCE CHARACTERISTICS:
LENGTH: 71
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: RNA
FEATURE:
OTHER INFORMATION: All pyrimidines are 2'-fluoro

US-08-870-930-25

Query Match 69.0%; Score 13.8; DB 4; Length 71;
Best Local Similarity 64.7%; Pred. No. 1.3e+02;
Matches 11; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGA 17
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Db 43 GAUUUCCUGCCGUCAGA 59

RESULT 15

US-09-254-968-28
Sequence 28, Application US/09254968
Patent No. 6426335
GENERAL INFORMATION:
APPLICANT: NEBOJSA JANJIC, LARRY GOLD, PAUL G. SCHMIDT, CHANDRA VARGESE,

MICHAEL WILLIS
TITLE OF INVENTION: VASCULAR ENDOTHELIAL GROWTH FACTOR (VEGF) NUCLEIC
ACID LIGAND COMPLEXES
NUMBER OF SEQUENCES: 139
CORRESPONDENCE ADDRESS:
ADDRESSEE: Swanson and Bratschun, L.L.C.
STREET: 8400 East Prentice Avenue, Suite #200
CITY: Denver
STATE: Colorado
COUNTRY: USA
ZIP: 80111
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 1.4 Mb storage
COMPUTER: IBM compatible
(C) OPERATING SYSTEM: MS-DOS
SOFTWARE: Word 7.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/254,968
FILING DATE: 13-Mar-2000
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US97/18944
FILING DATE: 17 OCTOBER 1997
APPLICATION NUMBER: 08/739,109
FILING DATE: 25 OCTOBER 1996
APPLICATION NUMBER: 08/870,930
FILING DATE: 6 JUNE 1997
APPLICATION NUMBER: 08/897,351
FILING DATE: 21 JULY 1997
ATTORNEY/AGENT INFORMATION:
NAME: Barry J. Swanson
REGISTRATION NUMBER: 33,215
REFERENCE/DOCKET NUMBER: NEX61C/PCT-US
TELECOMMUNICATION INFORMATION:
TELEPHONE: (303) 793-3333
TELEFAX: (303) 793-3433
INFORMATION FOR SEQ ID NO: 28:
SEQUENCE CHARACTERISTICS:
LENGTH: 71
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: RNA
FEATURE:
OTHER INFORMATION: All pyrimidines are 2'-fluoro (2'-F)
modified
SEQUENCE DESCRIPTION: SEQ ID NO: 28:
US-09-254-968-28
Query Match 69.0%; Score 13.8; DB 4; Length 71;
Best Local Similarity 64.7%; Pred. NO. 1.3e+02;
Matches 11; Conservative 4; Mismatches 2; Indels 0; Gaps 0;
QY 1 GATTGCTGACGTCAGA 17
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Db 43 GAUUUCCUGCCGUCAGA 59

Search completed: December 12, 2002, 01:41:42
Job time : 24.2464 secs

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GenCore version 5.1.3
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:06:16 ; Search time 318.116 Seconds
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Title: US-09-355-254F-13

Perfect score: 20
Sequence: 1 tgcagattgcgcaatctgca 20

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Gapop 10_0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0
Maximum DB seq length: 100

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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- 36: em_htg_mam.*
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- 38: em_sy.*
- 39: em_htgo_hum.*
- 40: em_htgo_mus.*
- 41: em_htgo_other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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4	20	100.0	20	6	A90879	A90879 Sequence 14
5	20	100.0	20	6	AR148641	AR148641 Sequence
6	20	100.0	20	6	AR148641	AR148641 Sequence
7	20	100.0	20	6	AX455640	AX455640 Sequence
8	20	100.0	20	6	AX455640	AX455640 Sequence
9	15.4	77.0	24	6	AX384713	AX384713 Sequence
10	15.4	77.0	24	6	AX384713	AX384713 Sequence
11	13.6	68.0	90	6	AR024336	AR024336 Sequence
12	13.6	68.0	90	6	AR024336	AR024336 Sequence
13	13.6	68.0	90	6	AR045189	AR045189 Sequence
14	13.6	68.0	90	6	AR045189	AR045189 Sequence
15	13.6	68.0	90	6	BD011413	BD011413 Chimeric
16	13.6	68.0	90	6	BD011413	BD011413 Chimeric
17	13.6	68.0	90	6	E43883	E43883 Chimeric an
18	13.6	68.0	90	6	E43883	E43883 Chimeric an
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20	13.4	67.0	28	6	AR202106	AR202106 Sequence
21	13.2	66.0	21	6	AR041200	AR041200 Sequence
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25	13.2	66.0	21	6	AR159952	AR159952 Sequence
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27	13.2	66.0	21	6	AX268106	AX268106 Sequence
28	13.2	66.0	33	6	AX268106	AX268106 Sequence
29	13.2	66.0	33	6	I65281	I65281 Sequence 3
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34	13.2	66.0	59	6	AX015212	AX015212 Sequence
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36	13.2	66.0	59	6	E64355	E64355 Single-stra
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41	12.8	64.0	56	6	AR077589	AR077589 Sequence
42	12.8	64.0	56	6	AR077589	AR077589 Sequence
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44	12.8	64.0	56	6	AX022408	AX022408 Sequence
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ALIGNMENTS

RESULT 1

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LOCUS A89792 20 bp DNA linear PAT 22-JAN-2000
DEFINITION Sequence 14 from Patent WO9832462.
ACCESSION A89792
VERSION A89792.1 GI:6738306
KEYWORDS
SOURCE unidentified.
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Lipford,G.B. and Heeg,K.
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
JOURNAL Patent: WO 9832462-A 14 30-JUL-1998;

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LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
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  /organism="unidentified"
  /db_xref="taxon:32644"
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ORIGIN
Query Match      100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCAGATTGCGCAATCTGCA 20
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Db 1 TGCAGATTGCGCAATCTGCA 20

RESULT 2
A89792/c
LOCUS      A89792      20 bp      DNA      linear      PAT 22-JAN-2000
DEFINITION Sequence 14 from Patent WO9832462.
ACCESSION  A89792
VERSION     A89792.1 GI:6738306
KEYWORDS   .
SOURCE     .
ORGANISM   unidentified.
            unclassified.
REFERENCE  1 (bases 1 to 20)
AUTHORS   Lipford,G.B. and Heeg,K.
TITLE     PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
            OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
JOURNAL    Patent: WO 9832462-A 14 30-JUL-1998;
            LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
FEATURES   Location/Qualifiers
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Best Local Similarity 100.0%; Pred. No. 22;
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A90879
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DEFINITION Sequence 14 from Patent EP0855184.
ACCESSION  A90879
VERSION     A90879.1 GI:6739288
KEYWORDS   .
SOURCE     .
ORGANISM   unidentified.
            unclassified.
REFERENCE  1 (bases 1 to 20)
AUTHORS   Heeg,K.P. and Lipford,G.B.
TITLE     PHARMACEUTICAL COMPOSITION COMPRISING A POLYNUCLEOTIDE AND AN
            ANTIGEN ESPECIALLY FOR VACCINATION
JOURNAL    Patent: EP 0855184-A 14 29-JUL-1998;
            HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
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Best Local Similarity 100.0%; Pred. No. 22;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 20 TGCAGATTGCGCAATCTGCA 1

RESULT 4
A90879/c
LOCUS      A90879      20 bp      DNA      linear      PAT 22-JAN-2000
DEFINITION Sequence 14 from Patent EP0855184.
ACCESSION  A90879
VERSION     A90879.1 GI:6739288
KEYWORDS   .
SOURCE     .
ORGANISM   unidentified.
            unclassified.
REFERENCE  1 (bases 1 to 20)
AUTHORS   Heeg,K.P. and Lipford,G.B.
TITLE     PHARMACEUTICAL COMPOSITION COMPRISING A POLYNUCLEOTIDE AND AN
            ANTIGEN ESPECIALLY FOR VACCINATION
JOURNAL    Patent: EP 0855184-A 14 29-JUL-1998;
            HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
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DEFINITION Sequence 7 from patent US 6225444.
ACCESSION  ARI48641
VERSION     ARI48641.1 GI:15112731
KEYWORDS   .
SOURCE     .
ORGANISM   Unknown.
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REFERENCE  1 (bases 1 to 20)
AUTHORS   Shashoua,V.E.
TITLE     Neuroprotective peptides and uses thereof
JOURNAL    Patent: US 6225444-A 7 01-MAY-2001;
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LOCUS      ARI48641      20 bp      DNA      linear      PAT 08-AUG-2001
DEFINITION Sequence 7 from patent US 6225444.

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ACCESSION ARI48641
VERSION ARI48641.1 GI:15112731
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Shashoua, V.E.
TITLE Neuroprotective peptides and uses thereof
JOURNAL Patent: US 6225444-A 7 01-MAY-2001;
FEATURES Location/Qualifiers
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RESULT 7
LOCUS AX455640 20 bp DNA linear PAT 06-JUL-2002
DEFINITION Sequence 117 from Patent WO0222809.
ACCESSION AX455640
VERSION AX455640.1 GI:21714708
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1
AUTHORS Bauer, S., Lipford, G. and Wagner, H.
TITLE Process for high throughput screening of cpq-based immuno-agonist/antagonist
JOURNAL Patent: WO 0222809-A 117 21-MAR-2002;
Coley Pharmaceutical GmbH (DE)
FEATURES Location/Qualifiers
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LOCUS AX455640/c 20 bp DNA linear PAT 06-JUL-2002
DEFINITION Sequence 117 from Patent WO0222809.
ACCESSION AX455640
VERSION AX455640.1 GI:21714708
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1
AUTHORS Bauer, S., Lipford, G. and Wagner, H.
TITLE Process for high throughput screening of cpq-based immuno-agonist/antagonist
JOURNAL Patent: WO 0222809-A 117 21-MAR-2002;
Coley Pharmaceutical GmbH (DE)
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DEFINITION Sequence 4 from Patent WO0181375.
ACCESSION AX384713
VERSION AX384713.1 GI:19577904
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1
AUTHORS Schepartz Shrader, A., Chin, J.W., Zutshi, R., Rutledge, S.E., Kehlbeck Martin, J.D. and Zondlo, N.J.
TITLE Dna & protein binding miniature proteins
JOURNAL Patent: WO 0181375-A 4 01-NOV-2001;
YALE UNIVERSITY (US)
FEATURES Location/Qualifiers
source 1..24
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DEFINITION Sequence 4 from Patent WO0181375.
ACCESSION AX384713
VERSION AX384713.1 GI:19577904
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1
AUTHORS Schepartz Shrader, A., Chin, J.W., Zutshi, R., Rutledge, S.E., Kehlbeck Martin, J.D. and Zondlo, N.J.
TITLE Dna & protein binding miniature proteins
JOURNAL Patent: WO 0181375-A 4 01-NOV-2001;
YALE UNIVERSITY (US)
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ORIGIN

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LOCUS AR024336 90 bp DNA linear PAT 05-DEC-1998
DEFINITION Sequence 104 from patent US 5795965.
ACCESSION AR024336
VERSION AR024336.1 GI:3977630
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 90)
AUTHORS Tsuchiya,M., Sato,K., Bendig,M.Margaret., Jones,S.Tarran. and Saldanha,J.William.
TITLE Reshaped human antibody to human interleukin-6 receptor
JOURNAL Patent: US 5795965-A 104 18-AUG-1998;
FEATURES Location/Qualifiers
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BASE COUNT 18 a 22 c 29 g 21 t
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LOCUS AR024336 90 bp DNA linear PAT 05-DEC-1998
DEFINITION Sequence 104 from patent US 5795965.
ACCESSION AR024336
VERSION AR024336.1 GI:3977630
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 90)
AUTHORS Tsuchiya,M., Sato,K., Bendig,M.Margaret., Jones,S.Tarran. and Saldanha,J.William.
TITLE Reshaped human antibody to human interleukin-6 receptor
JOURNAL Patent: US 5795965-A 104 18-AUG-1998;
FEATURES Location/Qualifiers
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BASE COUNT 18 a 22 c 29 g 21 t
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RESULT 13
AR045189

LOCUS AR045189 90 bp DNA linear PAT 29-SEP-1999
DEFINITION Sequence 104 from patent US 5817790.
ACCESSION AR045189
VERSION AR045189.1 GI:5966654
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 90)
AUTHORS Tsuchiya,M., Sato,K., Bendig,M.Margaret., Jones,S.Tarran. and Saldanha,J.William.
TITLE Reshaped human antibody to human interleukin-6 receptor
JOURNAL Patent: US 5817790-A 104 06-OCT-1998;
FEATURES Location/Qualifiers
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RESULT 14
AR045189/c
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DEFINITION Sequence 104 from patent US 5817790.
ACCESSION AR045189
VERSION AR045189.1 GI:5966654
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 90)
AUTHORS Tsuchiya,M., Sato,K., Bendig,M.Margaret., Jones,S.Tarran. and Saldanha,J.William.
TITLE Reshaped human antibody to human interleukin-6 receptor
JOURNAL Patent: US 5817790-A 104 06-OCT-1998;
FEATURES Location/Qualifiers
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BASE COUNT 18 a 22 c 29 g 21 t
ORIGIN

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Matches 16; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

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RESULT 15
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LOCUS BD011413 90 bp DNA linear PAT 31-JAN-2002
DEFINITION Chimeric antibody against human interleukin-6 receptor.
ACCESSION BD011413
VERSION BD011413.1 GI:18639786
KEYWORDS JP 2001083151-A/87.
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 90)
AUTHORS Tsuchiya,M., Sato,K., Bendig,M.M., Jones,S.T. and Saldanha,H.W.
TITLE Chimeric antibody against human interleukin-6 receptor
JOURNAL Patent: JP 2001083151-A 87 30-MAR-2001;
CHUGAI PHARMACEUTICAL CO LTD

COMMENT OS- Artificial Sequence
PN JP 2001083151-A/87
PD 30-MAR-2001
PF 28-JUL-2000 JP 2000229748
PR
PI MASAYUKI TSUCHIYA, KO SATO, MARY MARGARET BENDIGU, PI STEVEN
TAREN JONES,
PI HOSE WILLIAM SALDANHA
PC G01N33/53, A61K38/00, A61K39/395, A61K39/395, A61P35/00, PC
G01N33/577//C07K16/28,
PC C07K19/00, C12N15/09, (C12N15/09, C12R1:91), A61K37/02, C12N15/00,
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GenCore version 5.1.3
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Perfect score: 18
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Gapop 10.0 , Gapext 1.0

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SUMMARIES

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1	18	100.0	18	6	A89788	A89788 Sequence 10
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C 5	13.8	76.7	89	6	AX438437	AX438437 Sequence
C 6	13.2	73.3	82	9	HUMMACAJ	L37722 Homo sapien
C 7	13.2	73.3	88	10	MMALDRED04	U89144 Mus musculu
C 8	13	72.2	89	10	MMTCRAC2	X02846 M.musculus
9	12.8	71.1	29	6	AX099976	AX099976 Sequence
10	12.8	71.1	50	6	A16634	A16634 Nucleotide
11	12.8	71.1	50	6	A16645	A16645 Nucleotide
C 12	12.4	68.9	26	6	AX419077	AX419077 Sequence
C 13	12.4	68.9	51	6	AX204048	AX204048 Sequence
C 14	12.4	68.9	67	9	HSNVO7A55	U61467 Human mvosi
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16	12.4	68.9	85	6	AX360360	AX360360 Sequence
17	12.2	67.8	30	6	AX469615	AX469615 Sequence
18	12.2	67.8	50	6	AX165806	AX165806 Sequence
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21	12.2	67.8	89	6	I18551	I18551 Sequence 20
22	12.2	67.8	89	6	I34095	I34095 Sequence 20
C 23	12.2	67.8	94	1	ECWITA	X75467 E. coli wit
C 24	12.2	67.8	98	11	MMSTS6	Z36555 M.musculus
C 25	12.2	67.8	98	14	AF264011	AF264011 Hepatitis
C 26	12.2	67.8	100	5	XLRPS4	X64205 X.laavis mr
27	12.2	67.8	100	5	XLRPS11	X64208 X.laavis mr
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C 30	12	66.7	94	6	AX396657	AX396657 Sequence
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C 32	11.8	65.6	28	6	E38332	E38332 Process for
C 33	11.8	65.6	47	6	AX194676	AX194676 Sequence
C 34	11.8	65.6	52	6	E21658	E21658 Spermatogen
C 35	11.8	65.6	74	12	AF405700	AF405700 Synthetic
C 36	11.8	65.6	75	10	MUSIGHCA1	M31028 Mus musculu
C 37	11.8	65.6	98	6	AR081580	AR081580 Sequence
C 38	11.8	65.6	98	6	AX107888	AX107888 Sequence
39	11.6	64.4	25	6	AR089894	AR089894 Sequence
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C 41	11.6	64.4	25	6	AX116304	AX116304 Sequence
C 42	11.6	64.4	51	6	AX116305	AX116305 Sequence
43	11.6	64.4	51	6	AX158271	AX158271 Sequence
44	11.6	64.4	51	6	AX158272	AX158272 Sequence
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ALIGNMENTS

RESULT 1
A89788
LOCUS A89788 18 bp DNA linear PAT 22-JAN-2000
DEFINITION Sequence 10 from Patent WO9832462.
ACCESSION A89788
VERSION A89788.1 GI:6738302
KEYWORDS
SOURCE unidentified.
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 18)
AUTHORS Lipford G.B. and Heeg K.
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
JOURNAL Patent: WO 9832462-A 10 30-JUL-1998;

LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
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BASE COUNT 3 a 4 c 6 g 5 t
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LOCUS 18 bp DNA linear PAT 22-JAN-2000
DEFINITION Sequence 10 from Patent EP0855184.
A90875
ACCESSION A90875
VERSION A90875.1 GI:6739272
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 18)
AUTHORS Heeg,K.P. and Lipford,G.B.
TITLE Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination
JOURNAL Patent: EP 0855184-A 10 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
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RESULT 3
AX455583
LOCUS 18 bp DNA linear PAT 06-JUL-2002
DEFINITION Sequence 60 from Patent WO222809.
AX455583
ACCESSION AX455583
VERSION AX455583.1 GI:21714651
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1
AUTHORS Bauer, S., Lipford, G. and Wagner, H.
TITLE Process for high throughput screening of cpq-based immuno-agonist/antagonist
JOURNAL Patent: WO 0222809-A 60 21-MAR-2002;
Coley Pharmaceutical GmbH (DE)
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Source Location/Qualifiers
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BASE COUNT 3 a 4 c 6 g 5 t
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LOCUS 38 bp DNA linear PAT 07-OCT-1997
DEFINITION Sequence 10 from patent US 5648465.
I56082
ACCESSION I56082
VERSION I56082.1 GI:2476876
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 38)
AUTHORS Margolis,R.U., Rauch,U. and Margolis,R.K.
TITLE Cloning and expression of neurocan, a chondroitin sulfate proteoglycan
JOURNAL Patent: US 5648465-A 10 15-JUL-1997;
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QY 1 GGAATGACGTTCCCTGT 17
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Db 38 GGAATGATGTCNCNTGY 22

RESULT 5
AX438437/c
LOCUS 89 bp DNA linear PAT 28-JUN-2002
DEFINITION Sequence 6852 from Patent WO0229113.
AX438437
ACCESSION AX438437
VERSION AX438437.1 GI:21663245
KEYWORDS
SOURCE Bacillus clausii.
ORGANISM Bacillus clausii.
REFERENCE 1
AUTHORS Berka,R. and Clausen,I.G.
TITLE Methods for monitoring multiple gene expression
JOURNAL Patent: WO 0229113-A 6852 11-APR-2002;
Novozymes Biotech, Inc. (US); Novozymes A/S (DK)
FEATURES
Source Location/Qualifiers
1..89
/organism="Bacillus clausii"
/db_xref="taxon:79880"
21 a 16 c 29 g 23 t

BASE COUNT 21 a 16 c 29 g 23 t
ORIGIN

Query Match 76.7%; Score 13.8; DB 6; Length 89;
Best Local Similarity 88.2%; Pred. No. 6.5e+03;
Matches 15; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAATGACGTTCCCTGTG 18
|||||
Db 88 GAAGACATTCCTGTG 72

RESULT 6
HUMMACAJ/c
LOCUS 82 bp mRNA linear PRI 11-JAN-1995


```

DEFINITION Homo sapiens (clone 10) macronuclear mRNA.
ACCESSION L37722
VERSION L37722.1 GI:576851
KEYWORDS macronuclear.
SOURCE Homo sapiens cDNA to mRNA.
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 82)
AUTHORS Carney,J.P., McKnight,C.E., VanEpps,S. and Kelley,M.R.
TITLE Random amplification of cDNA ends (RRACE) allows for cloning of
multiple novel human cDNA fragments containing CAG repeats
JOURNAL Gene (1994) In press
COMMENT clones were isolated using a CAG oligo. The oligo was made up of
CAG x 8,
so by definition the first 24 bases are CAG x 8 (not included in
the
database entries). The actual number of CAG's is undetermined.
FEATURES
source
Location/Qualifiers
1..82
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="10"
/cell_line="Jurkat"
1..82
20 a 29 c 21 g 10 t 2 others
BASE COUNT 20 a 29 c 21 g 10 t 2 others
ORIGIN
Query Match 73.3%; Score 13.2; DB 9; Length 82;
Best Local Similarity 83.3%; Pred. No. 1.5e+04;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18
II IIIIIIIIIII
Db 64 GGCTGACGTTCCCTGTG 47

RESULT 7
MMALDRED04/c
LOCUS Mus musculus aldose reductase gene, exon 4.
DEFINITION 88 bp DNA linear ROD 13-APR-1998
ACCESSION U89144
VERSION U89144.1 GI:3046239
KEYWORDS
SEGMENT 4 of 10
SOURCE Mus musculus.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sclurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 88)
AUTHORS McGowan,M.H., Iwata,T. and Carper,D.A.
TITLE Characterization of the mouse aldose reductase gene and promoter in
a lens epithelial cell line
JOURNAL Mol. Vis. 4, 2 (1998)
MEDLINE 98153248
PUBMED 9485485
REMARK http://www.emory.edu/molvis/v4/p2
REFERENCE 2 (bases 1 to 88)
AUTHORS McGowan,M.H., Iwata,T. and Carper,D.A.
TITLE Direct Submission
JOURNAL Submitted (10-FEB-1997) LMOD, National Eye Institute, 9000
Rockville Pike, Bldg6/Rm232, Bethesda, MD 20892, USA
FEATURES
source
Location/Qualifiers
1..88
/organism="Mus musculus"
/strain="129 OLA"
/db_xref="taxon:10090"
/chromosome="6"
5..82
/number=4
21 a 23 c 24 g 20 t
BASE COUNT 21 a 23 c 24 g 20 t
ORIGIN

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Query Match 73.3%; Score 13.2; DB 10; Length 88;
Best Local Similarity 83.3%; Pred. No. 1.5e+04;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18
II IIIIIIIIIII
Db 51 GGTATCACGTTCCCTGAG 34

RESULT 8
MMTCRAC2
LOCUS M.musculus gene for Tcell receptor alpha-chain constant region exon
DEFINITION 2.
ACCESSION X02846
VERSION X02846.1 GI:54472
KEYWORDS constant region; T-cell receptor.
SOURCE Mus musculus.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sclurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 89)
AUTHORS Hayday,A.C., Diamond,D.J., Tanigawa,G., Heilig,J.S., Folsom,V.,
Saito,H. and Tonegawa,S.
TITLE Unusual organization and diversity of T-cell receptor alpha-chain
genes
JOURNAL Nature 316 (6031), 828-832 (1985)
MEDLINE 85296331
PUBMED 2993907
COMMENT The coding sequence is identical to that of alpha-chain mRNA
isolated from cytotoxic T-cells (clone PHDS58, see X01134). The
first base of the first codon of PHDS58 constant region is
identical to pos. 91 in X02843.
FEATURES
source
Location/Qualifiers
1..89
/organism="Mus musculus"
/db_xref="taxon:10090"
/cell_type="lymphocyte T"
<1..21
/note="intron"
22..66
/product="constant region of Tcell alpha chain"
/note="exon 2"
67..>89
/note="intron"
23 a 20 c 21 g 25 t
BASE COUNT 23 a 20 c 21 g 25 t
ORIGIN
Query Match 72.2%; Score 13; DB 10; Length 89;
Best Local Similarity 100.0%; Pred. No. 2e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 GAGGTTCCCTGTG 18
IIIIIIIIIIII
Db 21 GAGGTTCCCTGTG 33

RESULT 9
AX099976
LOCUS Sequence 5 from Patent WO0120007.
DEFINITION AX099976
ACCESSION AX099976
VERSION AX099976.1 GI:13538986
KEYWORDS synthetic construct.
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE 1 (bases 1 to 29)
AUTHORS Sillaots,S., Martinez-Perez,A., Tsang,A. and Storms,R.
TITLE A multifunctional system for the efficient manipulation of protein
expression in filamentous fungi and method using same
JOURNAL Patent: WO 0120007-A 5 22-MAR-2001;
Concordia University (CA)

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FEATURES
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        /organism="synthetic construct"
        /db_xref="taxon:32630"
        /note="Oligonucleotide"
BASE COUNT      8 a      8 c      5 g      8 t
ORIGIN

Query Match      71.1%; Score 12.8; DB 6; Length 29;
Best Local Similarity 87.5%; Pred. No. 2.8e+04;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY  2 GAATGACGTTCCCTG 17
    ||||| 11111
Db   2 GAATGAGTCCCTTT 17

RESULT 10
LOCUS      A16634      50 bp      DNA      linear      PAT 04-OCT-1994
DEFINITION Nucleotide sequence 9 from patent number EP0281530.
ACCESSION  A16634
VERSION     A16634.1 GI:641078
KEYWORDS   .
SOURCE     .
ORGANISM   .
REFERENCE  1
AUTHORS    Velati Bellini,A., Toma,S., Grandi,G., Colletti,E., Riboli,B.,
            Cosmina,P., Tognoni,A., Pedroni,P. and Rappuoli,R.
TITLE      Recombinant plasmids useable for the expression of heterologous
            proteins in bacillus
JOURNAL    Patent: EP 0281530-A 9 07-SEP-1988;
            ENRICERCHE S.p.A.; SCLAVO S.p.A
FEATURES   .
  source
    Location/Qualifiers
      1..50
        /organism="synthetic construct"
        /db_xref="taxon:32630"
BASE COUNT    11 a     12 c     15 g     12 t
ORIGIN

Query Match      71.1%; Score 12.8; DB 6; Length 50;
Best Local Similarity 87.5%; Pred. No. 2.7e+04;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY  1 GGAATGACGTTCCCTG 16
    ||||| 11111
Db   5 GGAATGACGATCCCTG 20

RESULT 11
LOCUS      A16645      50 bp      DNA      linear      PAT 04-OCT-1994
DEFINITION Nucleotide sequence 22 from patent number EP0281530.
ACCESSION  A16645
VERSION     A16645.1 GI:641088
KEYWORDS   .
SOURCE     .
ORGANISM   .
REFERENCE  1
AUTHORS    Velati Bellini,A., Toma,S., Grandi,G., Colletti,E., Riboli,B.,
            Cosmina,P., Tognoni,A., Pedroni,P. and Rappuoli,R.
TITLE      Recombinant plasmids useable for the expression of heterologous
            proteins in bacillus
JOURNAL    Patent: EP 0281530-A 22 07-SEP-1988;
            ENRICERCHE S.p.A.; SCLAVO S.p.A
FEATURES   .
  source
    Location/Qualifiers
      1..50
        /organism="synthetic construct"
        /db_xref="taxon:32630"
BASE COUNT    11 a     12 c     15 g     12 t
ORIGIN

Query Match      71.1%; Score 12.8; DB 6; Length 50;
Best Local Similarity 87.5%; Pred. No. 2.7e+04;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY  1 GGAATGACGTTCCCTG 16
    ||||| 11111
Db   5 GGAATGACGATCCCTG 20

RESULT 12
LOCUS      AX419077/c  26 bp      DNA      linear      PAT 18-JUN-2002
DEFINITION Sequence 96 from Patent WO0212471.
ACCESSION  AX419077
VERSION     AX419077.1 GI:21523851
KEYWORDS   .
SOURCE     .
ORGANISM   .
REFERENCE  1
AUTHORS    Acton,S., Robison,K.E. and Hsieh,P.Y.
TITLE      Angiotensin converting enzyme homolog and uses therefor
JOURNAL    Patent: WO 0212471-A 96 14-FEB-2002;
            Millennium Pharmaceuticals, Inc. (US)
FEATURES   .
  source
    Location/Qualifiers
      1..26
        /organism="synthetic construct"
        /db_xref="taxon:32630"
        /note="motifs"
BASE COUNT      7 a      3 c      9 g      7 t
ORIGIN

Query Match      68.9%; Score 12.4; DB 6; Length 26;
Best Local Similarity 92.9%; Pred. No. 4.8e+04;
Matches 13; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY  2 GAATGACGTTCCCT 15
    ||||| 11111
Db   16 GAATGACTTCCCT 3

RESULT 13
LOCUS      AX204048      51 bp      DNA      linear      PAT 30-AUG-2001
DEFINITION Sequence 154 from Patent WO0148245.
ACCESSION  AX204048
VERSION     AX204048.1 GI:15393526
KEYWORDS   .
SOURCE     .
ORGANISM   human.
REFERENCE  1 (bases 1 to 51)
AUTHORS    Shimkets,R.A. and Leach,M.
TITLE      Nucleic acids containing single nucleotide polymorphisms and
            methods of use thereof
JOURNAL    Patent: WO 0148245-A 154 05-JUL-2001;
            Curagen Corporation (US)
FEATURES   .
  source
    Location/Qualifiers
      1..51
        /organism="Homo sapiens"
        /db_xref="taxon:9606"
        /note="single nucleotide polymorphism"
BASE COUNT     16 a     15 c     13 g      7 t
ORIGIN

Query Match      68.9%; Score 12.4; DB 6; Length 51;
Best Local Similarity 92.9%; Pred. No. 4.6e+04;
Matches 13; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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QY 5 TGACGTTCCCTGTG 18
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Db 39 TGATGTTCCCTGTG 26

RESULT 14
HSMY07A55
LOCUS HSMY07A55 67 bp DNA linear PRI 19-MAR-1997
DEFINITION Human myosin VIIa (MYO7A) gene, 5' exon 38.
ACCESSION U61467
VERSION U61467.1 GI:1894878
KEYWORDS
SEGMENT
SOURCE
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 67)
AUTHORS Kelley.P.M., Weston.M.D., Chen.Z.-Y., Orten.D.J., Hasson.T.,
Overbeck.L.D., Pinniti.J., Talmadge.C.B., Ing.P., Mooseker.M.S.,
Corey.D., Sumegi.J. and Kimberling.W.J.
TITLE The genomic structure of the gene defective in Usher syndrome type
Ib (MYO7A)
JOURNAL Genomics 40 (1), 73-79 (1997)
MEDLINE 97224487
PUBMED 9070921
REFERENCE 2 (bases 1 to 67)
AUTHORS Kelley.P.M.
TITLE Direct Submission
JOURNAL Submitted (21-JUN-1996) Gene Marker Lab, Boys Town National
Research Hospital, 555 North 30th Street, Omaha, NE 68131, USA
FEATURES
source
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/organism="Homo sapiens"
/db_xref="taxon:9606"
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/gene="MYO7A"
/number=38
BASE COUNT 11 a 24 c 21 g 11 t
ORIGIN

Query Match 68.9%; Score 12.4; DB 9; Length 67;
Best Local Similarity 92.9%; Pred. No. 4.5e+04;
Matches 13; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 5 TGACGTTCCCTGTG 18
|||||
Db 19 TGACGTTCCCTGTG 32

RESULT 15
AR092212
LOCUS AR092212 70 bp DNA linear PAT 08-SEP-2000
DEFINITION Sequence 150 from patent US 598142.
ACCESSION AR092212
VERSION AR092212.1 GI:10018966
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 70)
AUTHORS Gold,L., Eaton,B., Smith,D., Wecker,M. and Jensen,K.
TITLE Systematic evolution of ligands by exponential enrichment:
chemi-SELEX
JOURNAL Patent: US 598142-A 150 07-DEC-1999;
FEATURES
source
1..70
/organism="unknown"
BASE COUNT 16 a 17 c 25 g 12 t
ORIGIN

Query Match 68.9%; Score 12.4; DB 6; Length 70;
Best Local Similarity 92.9%; Pred. No. 4.5e+04;

Matches 13; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 3 AATGACGTTCCCTG 16
|||||
Db 34 AATGACGTACCCTG 47

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Job time : 295.304 secs

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SUMMARIES

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Searched: 2054640 seqs, 14551402878 residues

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Listing first 45 summaries

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3	20	100.0	20	6	AX455587 Sequence
4	17.4	87.0	22	6	AX040434 Sequence
5	17	85.0	19	6	I81948 Sequence 46
6	15.4	77.0	17	6	I87792 Sequence 20
7	15.4	77.0	19	6	I81949 Sequence 47
8	15.4	77.0	22	6	AX040435 Sequence
9	15.4	77.0	25	6	AR182575 Sequence
10	15.4	77.0	25	6	AX472568 Sequence
11	15.4	77.0	25	6	AX476844 Sequence
12	15.4	77.0	25	6	AX476862 Sequence
13	15.4	77.0	28	6	I39727 Sequence 14
14	15.4	77.0	28	6	I55844 Sequence 14
15	15.2	76.0	20	6	AR103469 Sequence
16	15.2	76.0	20	6	AR176473 Sequence
17	14.4	72.0	18	6	AR153598 Sequence
18	14.2	71.0	26	6	AR160395 Sequence
19	14.2	71.0	26	6	AX482621 Sequence
20	14.2	71.0	72	6	I95007 Sequence 24
21	14.2	71.0	76	6	I95006 Sequence 23
22	14	70.0	19	6	AX338646 Sequence
23	13.8	69.0	25	6	AX192408 Sequence
24	13.8	69.0	28	6	I39716 Sequence 3
25	13.8	69.0	28	6	I55833 Sequence 3
26	13.6	68.0	35	6	AR061506 Sequence
27	13.6	68.0	35	6	AR108405 Sequence
28	13.6	68.0	35	6	I16362 Sequence 18
29	13.6	68.0	35	6	I66848 Sequence 18
30	13.6	68.0	35	6	I84942 Sequence 18
31	13.6	68.0	94	11	AL773118 Arabidops
32	13.2	66.0	66	10	MUSMUP
33	13.2	66.0	83	3	DME428807
34	13	65.0	13	6	AX026536 Sequence
35	12.8	64.0	52	6	AR103470 Sequence
36	12.8	64.0	52	6	AR176474 Sequence
37	12.6	63.0	22	6	AR062163 Sequence
38	12.6	63.0	30	6	AR004706 Sequence
39	12.6	63.0	30	6	AR008192 Sequence
40	12.6	63.0	30	6	AR136975 Sequence
41	12.6	63.0	30	6	I76976 Sequence 36
42	12.6	63.0	30	6	I80971 Sequence 36
43	12.6	63.0	30	6	I81067 Sequence 36
44	12.6	63.0	46	6	AR126436 Sequence
45	12.6	63.0	50	6	AR135509 Sequence

ALIGNMENTS

RESULT 1
A89795
LOCUS A89795 20 bp DNA linear PAT 22-JAN-2000
DEFINITION Sequence 17 from Patent WO9832462.
ACCESSION A89795
VERSION A89795.1 GI:6738309
KEYWORDS
SOURCE unidentified.
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Lipford,G.B. and Heeg,K.
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
JOURNAL Patent: WO 9832462-A 17 30-JUL-1998;

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FEATURES
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LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
Location/Qualifiers
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ORIGIN

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Best Local Similarity 100.0%; Pred. No. 6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCCATTTCCTCGTAATCTT 20
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Db 1 GTCCATTTCCTCGTAATCTT 20

RESULT 2
AX455587
LOCUS
DEFINITION
Sequence 64 from Patent WO222809.
ACCESSION
AX455587.1
VERSION
AX455587.1 GI:21714655
KEYWORDS
synthetic construct.
SOURCE
artificial sequences.
ORGANISM
Bauer, S., Lipford, G. and Wagner, H.
REFERENCE
1 (bases 1 to 20)
AUTHORS
Heeg, K.P. and Lipford, G.B.
TITLE
Pharmaceutical composition comprising a polynucleotide and an
antigen especially for vaccination
JOURNAL
Patent: EP 0855184-A 17 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
FEATURES
Location/Qualifiers
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/organism="unidentified"
/db_xref="taxon:32644"
BASE COUNT      4 a      6 c      2 g      8 t
ORIGIN

Query Match      100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCCATTTCCTCGTAATCTT 20
    |||||
Db 1 GTCCATTTCCTCGTAATCTT 20

RESULT 3
AX455587
LOCUS
DEFINITION
Sequence 64 from Patent WO222809.
ACCESSION
AX455587
VERSION
AX455587.1 GI:21714655
KEYWORDS
synthetic construct.
SOURCE
artificial sequences.
ORGANISM
Bauer, S., Lipford, G. and Wagner, H.
REFERENCE
1 (bases 1 to 20)
AUTHORS
Heeg, K.P. and Lipford, G.B.
TITLE
Pharmaceutical composition comprising a polynucleotide and an
antigen especially for vaccination
JOURNAL
Patent: EP 0855184-A 17 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
FEATURES
Location/Qualifiers
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/db_xref="taxon:32644"
BASE COUNT      4 a      6 c      2 g      8 t
ORIGIN

Query Match      100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCCATTTCCTCGTAATCTT 20
    |||||
Db 1 GTCCATTTCCTCGTAATCTT 20

RESULT 4
AX404034
LOCUS
DEFINITION
Sequence 9 from Patent WO063357.
ACCESSION
AX404034
VERSION
AX404034.1 GI:11230241
KEYWORDS
synthetic construct.
SOURCE
artificial sequences.
ORGANISM
Flier, J.S. and Bjorbaek, C.
REFERENCE
1 (bases 1 to 22)
AUTHORS
Methods and compositions for modulating ciliary neurotrophic facto
r activity
JOURNAL
Patent: WO 0063357-A 9 26-OCT-2000;
Beth Israel Deaconess Medical Center (US)
FEATURES
Location/Qualifiers
1..22
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="primer-bind"
BASE COUNT      5 a      8 c      2 g      7 t
ORIGIN

Query Match      87.0%; Score 17.4; DB 6; Length 22;
Best Local Similarity 94.7%; Pred. No. 1.5e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 TCCATTTCCTCGTAATCTT 20
    |||||
Db 4 TCCATTTCCTCGTAATCAT 22

RESULT 5
I81948
LOCUS
DEFINITION
Sequence 46 from patent US 5712094.
ACCESSION
I81948
VERSION
I81948.1 GI:3210245
KEYWORDS
Unknown.
SOURCE
Unclassified.
ORGANISM
Seidel, H. Martin., Lamb, I. Peter., and Chan, S.-S. Tian.
REFERENCE
1 (bases 1 to 19)
AUTHORS
Methods for detecting modulators of cytokine action
TITLE
Patent: US 5712094-A 46 27-JAN-1998;
JOURNAL
Location/Qualifiers
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FEATURES
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BASE COUNT      5 a      6 c      2 g      6 t
ORIGIN

Query Match      85.0%; Score 17; DB 6; Length 19;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 TCCATTTCCTCGTAATCTT 18
    |||||
Db 3 TCCATTTCCTCGTAATCTT 19

RESULT 6
I87792

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FEATURES	source	Location/Qualifiers
BASE COUNT.	6 a 4 c 6 g 6 t	
ORIGIN		
Query Match	77.0%; Score 15.4; DB 6; Length 22;	
Best Local Similarity	94.1%; Pred. No. 1.7e+03;	
Matches	16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;	
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DB		
22 CATTTCCTCGTAATCAT 6		
RESULT 9		
LOCUS	AR182575	25 bp DNA linear PAT 20-APR-2002
DEFINITION	Sequence 23 from patent US 6338949.	
ACCESSION	AR182575	
VERSION	AR182575.1 GI:20225782	
KEYWORDS	Unknown.	
SOURCE	Unknown.	
ORGANISM	Unclassified.	
REFERENCE	1 (bases 1 to 25)	
AUTHORS	Darnell,J.E. Jr., Schindler,C.W., Fu,X.-Y., Wen,Z. and Zhong,Z.	
TITLE	Nucleic acids encoding receptor recognition factor stat4 and methods of use thereof	
JOURNAL	Patent: US 6338949-A 23 15-JAN-2002;	
FEATURES	Location/Qualifiers	
source	1..25	
BASE COUNT	6 a 7 c 5 g 7 t	
ORIGIN	/organism="unknown"	
Query Match	77.0%; Score 15.4; DB 6; Length 25;	
Best Local Similarity	94.1%; Pred. No. 1.7e+03;	
Matches	16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;	
QY	4 CATTTCCTCGTAATCTTT 20	
DB		
6 CATTTCCTCGTAATCGT 22		
RESULT 10		
LOCUS	AX472568	25 bp DNA linear PAT 09-AUG-2002
DEFINITION	Sequence 63 from Patent WO02052039.	
ACCESSION	AX472568	
VERSION	AX472568.1 GI:22207472	
KEYWORDS	synthetic construct.	
SOURCE	synthetic construct	
ORGANISM	artificial sequences.	
REFERENCE	1	
AUTHORS	Blais,Y., Rousseau,P., Leblanc,B. and Camato,R.N.	
TITLE	Methods for selecting and producing selective pharmaceutical compounds and compositions using an established genetically altered cell-based library responsive to transcription factors; genetic constructs and library therefor	
JOURNAL	Patent: WO 02052039-A 63 04-JUL-2002;	
FEATURES	Geneka Biotechnology Inc. (CA)	
source	Location/Qualifiers	
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6 a 7 c 5 g 7 t	/db_xref="taxon:32630"	
	/note="Oligonucleotide"	

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 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CATTTCCCGTAATCTT 20
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 Db 6 CATTTCCCGTAATCTT 22

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 AX476844
 LOCUS AX476844 25 bp DNA linear PAT 12-AUG-2002
 DEFINITION Sequence 21 from Patent WO02052037.
 ACCESSION AX476844
 VERSION AX476844.1 GI:22216120
 KEYWORDS synthetic construct.
 SOURCE synthetic construct
 ORGANISM artificial sequences.
 REFERENCE 1
 AUTHORS Larose,A.M., Rousseau,P., Leblanc,B. and Camato,R.
 TITLE Method for screening and/or identifying factors that bind to nucleic acids
 JOURNAL Patent: WO 02052037-A 21 04-JUL-2002;
 FEATURES Location/Qualifiers
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 /organism="synthetic construct"
 /db_xref="taxon:32630"
 /note="NABE-probes"

BASE COUNT 6 a 7 c 5 g 7 t
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Query Match 77.0%; Score 15.4; DB 6; Length 25;
 Best Local Similarity 94.1%; Pred. No. 1.7e+03;
 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CATTTCCCGTAATCTT 20
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 Db 6 CATTTCCCGTAATCTT 22

RESULT 12
 AX476862
 LOCUS AX476862 25 bp DNA linear PAT 12-AUG-2002
 DEFINITION Sequence 39 from Patent WO02052037.
 ACCESSION AX476862
 VERSION AX476862.1 GI:22216138
 KEYWORDS synthetic construct.
 SOURCE synthetic construct
 ORGANISM artificial sequences.
 REFERENCE 1
 AUTHORS Larose,A.M., Rousseau,P., Leblanc,B. and Camato,R.
 TITLE Method for screening and/or identifying factors that bind to nucleic acids
 JOURNAL Patent: WO 02052037-A 39 04-JUL-2002;
 FEATURES Location/Qualifiers
 source 1..25
 /organism="synthetic construct"
 /db_xref="taxon:32630"
 /note="Double stranded NABE"

BASE COUNT 6 a 7 c 5 g 7 t
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 Best Local Similarity 94.1%; Pred. No. 1.7e+03;
 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CATTTCCCGTAATCTT 20
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 Db 6 CATTTCCCGTAATCTT 22

RESULT 13
 I39727
 LOCUS I39727 28 bp DNA linear PAT 13-MAY-1997
 DEFINITION Sequence 14 from patent US 5616489.
 ACCESSION I39727
 VERSION I39727.1 GI:2084207
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.

REFERENCE 1 (bases 1 to 28)
 AUTHORS Levy,D.E.
 TITLE DNA sequence which binds transcriptional regulatory proteins activated in response to various cytokines and uses thereof
 JOURNAL Patent: US 5616489-A 14 01-APR-1997;
 FEATURES Location/Qualifiers
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BASE COUNT 7 a 9 c 3 g 9 t
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 Best Local Similarity 94.1%; Pred. No. 1.7e+03;
 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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 Db 8 TTCATTTCCTCGTAATC 24

RESULT 14
 I55844
 LOCUS I55844 28 bp DNA linear PAT 07-OCT-1997
 DEFINITION Sequence 14 from patent US 5648217.
 ACCESSION I55844
 VERSION I55844.1 GI:2476638
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.

REFERENCE 1 (bases 1 to 28)
 AUTHORS Levy,D.E.
 TITLE DNA sequence which binds transcriptional regulatory proteins activated in response to various cytokines and uses thereof
 JOURNAL Patent: US 5648217-A 14 15-JUL-1997;
 FEATURES Location/Qualifiers
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BASE COUNT 7 a 9 c 3 g 9 t
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 Best Local Similarity 94.1%; Pred. No. 1.7e+03;
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QY 2 TCCATTTCCTCGTAATC 18
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 Db 8 TTCATTTCCTCGTAATC 24

RESULT 15
 AR103469
 LOCUS AR103469 AR103469 20 bp DNA linear PAT 14-FEB-2001
 DEFINITION Sequence 5 from patent US 6087478.
 ACCESSION AR103469
 VERSION AR103469.1 GI:12815057
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 20)

AUTHORS Vinkemeier, J., Moarefi, I., Darnell, J.E. Jr. and Kuriyan, J.
 TITLE Crystal of the N-terminal domain of a STAT protein and methods of
 use thereof
 JOURNAL Patent: US 6087478-A 5 11-JUL-2000;
 FEATURES Location/Qualifiers
 source 1..20
 BASE COUNT 5 a 6 c 3 g 6 t
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 Best Local Similarity 85.0%; Pred. No. 2.1e+03;
 Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
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 Search completed: December 12, 2002, 02:55:59
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Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues
Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0
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Post-processing: Minimum Match 0%
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Listing first 45 summaries

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- 3: gb_in.*
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- 41: em_hgtgo_other.*

Pred. No. is the number of results predicted by chance to have a

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and is derived by analysis of the total score distribution.

SUMMARIES

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2	20	100.0	20	6	A90883	A90883 Sequence 18
3	20	100.0	20	6	AX455591	AX455591 Sequence
4	19	95.0	24	6	AX406780	AX406780 Sequence
5	16	80.0	19	6	I79489	I79489 Sequence 75
6	16	80.0	19	6	I81942	I81942 Sequence 40
7	15	75.0	17	6	I87793	I87793 Sequence 21
8	15	75.0	19	6	I79490	I79490 Sequence 76
9	15	75.0	19	6	I81943	I81943 Sequence 41
10	14.8	74.0	47	6	AR032425	AR032425 Sequence
11	14.8	74.0	47	6	AR209089	AR209089 Sequence
12	14.8	74.0	47	6	I29165	I29165 Sequence 37
13	14.8	74.0	47	6	I90839	I90839 Sequence 37
14	14.8	74.0	50	6	AR032824	AR032824 Sequence
15	14.8	74.0	50	6	AR032825	AR032825 Sequence
16	14.8	74.0	50	6	AR209488	AR209488 Sequence
17	14.8	74.0	50	6	AR209489	AR209489 Sequence
18	14.8	74.0	50	6	I29564	I29564 Sequence 43
19	14.8	74.0	50	6	I29565	I29565 Sequence 43
20	14.8	74.0	50	6	I91238	I91238 Sequence 43
21	14.8	74.0	50	6	I91239	I91239 Sequence 43
22	14.4	72.0	19	6	I79437	I79437 Sequence 23
23	14.4	72.0	19	6	I79439	I79439 Sequence 23
24	14.4	72.0	19	6	I79447	I79447 Sequence 33
25	14.4	72.0	19	6	I79463	I79463 Sequence 49
26	14.4	72.0	19	6	I79465	I79465 Sequence 51
27	14.4	72.0	19	6	I79469	I79469 Sequence 55
28	14.4	72.0	19	6	I81944	I81944 Sequence 42
29	14.4	72.0	32	6	E09298	E09298 DNA linker
30	13.6	68.0	50	6	E40770	E40770 Antihuman F
31	13.4	67.0	19	6	I79438	I79438 Sequence 24
32	13.4	67.0	19	6	I79440	I79440 Sequence 26
33	13.4	67.0	19	6	I79448	I79448 Sequence 34
34	13.4	67.0	19	6	I79464	I79464 Sequence 50
35	13.4	67.0	19	6	I79466	I79466 Sequence 52
36	13.4	67.0	19	6	I79470	I79470 Sequence 56
37	13.4	67.0	19	6	I81945	I81945 Sequence 43
38	13.4	67.0	24	6	AX444661	AX444661 Sequence
39	13.4	67.0	25	6	AX116799	AX116799 Sequence
40	13.4	67.0	51	6	AX163145	AX163145 Sequence
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ALIGNMENTS

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LOCUS A89796
DEFINITION Sequence 18 from Patent WO9832462.
ACCESSION A89796
VERSION A89796.1 GI:6738310
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Lipford, G. B. and Heeg, K.
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
JOURNAL Patent: WO 9832462-A 18 30-JUL-1998;
linear PAT 22-JAN-2000

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DEFINITION Sequence 68 from Patent WO0222809.
ACCESSION AX455591
VERSION AX455591.1 GI:21714659
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequences.
REFERENCE
AUTHORS Bauer, S., Lipford, G. and Wagner, H.
TITLE Process for high throughput screening of cpg-based
JOURNAL immuno-agonist/antagonist
PATENT: WO 0222809-A 68 21-MAR-2002;
Cole Pharmaceutical GmbH (DE)
FEATURES
Location/Qualifiers
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/db_xref="taxon:32630"
/Note="Synthetic oligonucleotide"
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BASE COUNT
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RESULT 3
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LOCUS
DEFINITION Sequence 68 from Patent WO0222809.
ACCESSION AX455591
VERSION AX455591.1 GI:21714659
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequences.
REFERENCE
AUTHORS Bauer, S., Lipford, G. and Wagner, H.
TITLE Process for high throughput screening of cpg-based
JOURNAL immuno-agonist/antagonist
PATENT: WO 0222809-A 68 21-MAR-2002;
Cole Pharmaceutical GmbH (DE)
FEATURES
Location/Qualifiers
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/db_xref="taxon:32630"
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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DEFINITION Sequence 36 from Patent WO0229044.
ACCESSION AX406780
VERSION AX406780.1 GI:21439705
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequences.
REFERENCE
AUTHORS Hecker, M. and Wagner, A.H.
TITLE Modulation of the transcription of pro-inflammatory gene products
JOURNAL Patent: WO 0229044-A 36 11-APR-2002;
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Location/Qualifiers
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/organism="synthetic construct"
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/Note="Oligonucleotide"
6 a 4 c 4 g 10 t

BASE COUNT
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ORIGIN

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Db 6 TATGCATATTCCTGTAAGT 24

RESULT 5
I79489
LOCUS
DEFINITION Sequence 75 from patent US 5707803.
ACCESSION I79489
VERSION I79489.1 GI:3207779
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE
AUTHORS Lamb, I. Peter. and Seidel, H. Martin.
TITLE DNA regulatory elements responsive to cytokines and methods for
JOURNAL their use
PATENT: US 5707803-A 75 13-JAN-1998;
FEATURES
Location/Qualifiers
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5 a 3 c 4 g 7 t

BASE COUNT
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ORIGIN

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Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 5 CATATTCTCTGTAAGTG 20
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Db 4 CATATTCTCTGTAAGTG 19

RESULT 6
I81942
LOCUS

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DEFINITION Sequence 40 from patent US 5712094.
ACCESSION I81942
VERSION I81942.1 GI:3210239
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 19)
AUTHORS Seidel, H. Martin., Lamb, I. Peter. and Chan, S.-S. Tian.
TITLE Methods for detecting modulators of cytokine action
JOURNAL Patent: US 5712094-A 40 27-JAN-1998;
FEATURES Location/Qualifiers
source 1..19
BASE COUNT 5 a 3 c 4 g 7 t
ORIGIN

Query Match 80.0%; Score 16; DB 6; Length 19;
Best Local Similarity 100.0%; Pred. No. 3.2e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 4 CATATTCCTGTAAGTG 19

RESULT 7
I87793
LOCUS
DEFINITION Sequence 21 from patent US 5716622.
ACCESSION I87793
VERSION I87793.1 GI:3407733
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 17)
AUTHORS Darnell, J.E. Jr., Wen, Z., Horvath, C.M. and Zhong, Z.
TITLE Functionally active regions of signal transducer and activators of transcription
JOURNAL Patent: US 5716622-A 21 10-FEB-1998;
FEATURES Location/Qualifiers
source 1..17
BASE COUNT 5 a 2 c 3 g 7 t
ORIGIN

Query Match 75.0%; Score 15; DB 6; Length 17;
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Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATTCCTGTAAGTG 20
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Db 1 ATATTCCTGTAAGTG 15

RESULT 8
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LOCUS
DEFINITION Sequence 76 from patent US 5707803.
ACCESSION I79490
VERSION I79490.1 GI:3207780
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 19)
AUTHORS Lamb, I. Peter. and Seidel, H. Martin.
TITLE DNA regulatory elements responsive to cytokines and methods for their use
JOURNAL Patent: US 5707803-A 76 13-JAN-1998;
FEATURES Location/Qualifiers
source 1..19

BASE COUNT 7 a 4 c 3 g 5 t
ORIGIN

Query Match 75.0%; Score 15; DB 6; Length 19;
Best Local Similarity 100.0%; Pred. No. 9.9e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATTCCTGTAAGTG 20
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I81943/c
LOCUS
DEFINITION Sequence 41 from patent US 5712094.
ACCESSION I81943
VERSION I81943.1 GI:3210240
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 19)
AUTHORS Seidel, H. Martin., Lamb, I. Peter. and Chan, S.-S. Tian.
TITLE Methods for detecting modulators of cytokine action
JOURNAL Patent: US 5712094-A 41 27-JAN-1998;
FEATURES Location/Qualifiers
source 1..19
BASE COUNT 7 a 4 c 3 g 5 t
ORIGIN

Query Match 75.0%; Score 15; DB 6; Length 19;
Best Local Similarity 100.0%; Pred. No. 9.9e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATTCCTGTAAGTG 20
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Db 19 ATATTCCTGTAAGTG 5

RESULT 10
AR032425/c
LOCUS
DEFINITION Sequence 37 from patent US 5869241.
ACCESSION AR032425
VERSION AR032425.1 GI:5948030
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 47)
AUTHORS Edwards, C.A., Cantor, C.R., Andrews, B.M., Turin, L.M. and Fry, K.E.
TITLE Method of determining DNA sequence preference of a DNA-binding molecule
JOURNAL Patent: US 5869241-A 37 09-FEB-1999;
FEATURES Location/Qualifiers
source 1..47
BASE COUNT 17 a 6 c 10 g 14 t
ORIGIN

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Db 28 TATTATATTCCTGTAAG 11

RESULT 11

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LOCUS AR209089 47 bp DNA linear PAT 20-JUN-2002
DEFINITION Sequence 37 from patent US 6384208.
ACCESSION AR209089
VERSION AR209089.1 GI:21510414
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE
1 (bases 1 to 47)
Unclassified.
AUTHORS Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.
TITLE Sequence directed DNA binding molecules compositions and methods
JOURNAL Patent: US 6384208-A 37 07-MAY-2002;
FEATURES
Location/Qualifiers
1..47
Source

BASE COUNT 17 a 6 c 10 g 14 t
ORIGIN

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Best Local Similarity 88.9%; Pred. No. 1.1e+04;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18
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Db 28 TATTTATATTCCTGTAAG 11

RESULT 12
LOCUS I29165/c 47 bp DNA linear PAT 06-FEB-1997
DEFINITION Sequence 37 from patent US 5578444.
ACCESSION I29165
VERSION I29165.1 GI:1819956
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE
1 (bases 1 to 47)
Unclassified.
AUTHORS Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.
TITLE Sequence-directed DNA-binding molecules compositions and methods
JOURNAL Patent: US 5578444-A 37 26-NOV-1996;
FEATURES
Location/Qualifiers
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Source

BASE COUNT 17 a 6 c 10 g 14 t
ORIGIN

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Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Db 28 TATTTATATTCCTGTAAG 11

RESULT 13
LOCUS I90839/c 47 bp DNA linear PAT 01-DEC-1998
DEFINITION Sequence 37 from patent US 5726014.
ACCESSION I90839
VERSION I90839.1 GI:3935309
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE
1 (bases 1 to 47)
Unclassified.
AUTHORS Edwards,C.A., Cantor,C.R., Andrews,B.M. and Turin,L.M.
TITLE Screening assay for the detection of DNA-binding molecules
JOURNAL Patent: US 5726014-A 37 10-MAR-1998;
FEATURES
Location/Qualifiers
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Source

BASE COUNT 17 a 6 c 10 g 14 t
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Query Match 74.0%; Score 14.8; DB 6; Length 47;
Best Local Similarity 88.9%; Pred. No. 1.1e+04;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18
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Db 28 TATTTATATTCCTGTAAG 11

RESULT 14
LOCUS AR032824/c 50 bp DNA linear PAT 29-SEP-1999
DEFINITION Sequence 436 from patent US 5869241.
ACCESSION AR032824
VERSION AR032824.1 GI:5948429
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE
1 (bases 1 to 50)
Unclassified.
AUTHORS Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.
TITLE Method of determining DNA sequence preference of a DNA-binding molecule.
JOURNAL Patent: US 5869241-A 436 09-FEB-1999;
FEATURES
Location/Qualifiers
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Source

BASE COUNT 18 a 6 c 10 g 16 t
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RESULT 15
LOCUS AR032825/c 50 bp DNA linear PAT 29-SEP-1999
DEFINITION Sequence 437 from patent US 5869241.
ACCESSION AR032825
VERSION AR032825.1 GI:5948430
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE
1 (bases 1 to 50)
Unclassified.
AUTHORS Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.
TITLE Method of determining DNA sequence preference of a DNA-binding molecule.
JOURNAL Patent: US 5869241-A 437 09-FEB-1999;
FEATURES
Location/Qualifiers
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BASE COUNT 18 a 6 c 10 g 16 t
ORIGIN

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Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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GenCore version 5.1.3
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1829.698 Million cell updates/sec

Title: US-09-355-254F-22

Perfect score: 20
Sequence: 1 aagcgaatgaaattgact 20

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0
Maximum DB seq length: 100

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

GenEmbl: *

- 1: gb_ba:*
- 2: gb_htg:*
- 3: gb_in:*
- 4: gb_om:*
- 5: gb_ov:*
- 6: gb_pat:*
- 7: gb_ph:*
- 8: gb_pl:*
- 9: gb_pr:*
- 10: gb_ro:*
- 11: gb_sts:*
- 12: gb_sy:*
- 13: gb_un:*
- 14: gb_vi:*
- 15: em_ba:*
- 16: em_fun:*
- 17: em_hum:*
- 18: em_in:*
- 19: em_mu:*
- 20: em_om:*
- 21: em_or:*
- 22: em_ov:*
- 23: em_pat:*
- 24: em_ph:*
- 25: em_pl:*
- 26: em_ro:*
- 27: em_sts:*
- 28: em_un:*
- 29: em_vi:*
- 30: em_htg_hum:*
- 31: em_htg_inv:*
- 32: em_htg_other:*
- 33: em_htg_mus:*
- 34: em_htg_pln:*
- 35: em_htg_rod:*
- 36: em_htg_mam:*
- 37: em_htg_vrt:*
- 38: em_sy:*
- 39: em_htgo_hum:*
- 40: em_htgo_mus:*
- 41: em_htgo_other:*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	20	100.0	20	6	A89801	A89801 Sequence 23
2	20	100.0	20	6	A90888	A90888 Sequence 23
3	20	100.0	20	6	AX455545	AX455545 Sequence
4	20	100.0	22	6	AX406763	AX406763 Sequence
5	20	100.0	22	6	AX406764	AX406764 Sequence
6	14.8	74.0	65	6	AX482985	AX482985 Sequence
7	14.4	72.0	21	6	AX392034	AX392034 Sequence
8	13.8	69.0	20	6	AX162873	AX162873 Sequence
9	13.8	69.0	20	6	AR194023	AR194023 Sequence
10	13.8	69.0	51	6	AX204090	AX204090 Sequence
11	13.6	68.0	24	6	AR174147	AR174147 Sequence
12	13.6	68.0	24	6	AX456503	AX456503 Sequence
13	13.6	68.0	50	6	AX060077	AX060077 Sequence
14	13.6	68.0	51	6	AX157392	AX157392 Sequence
15	13.6	68.0	89	9	HUMADPRT11	M29774 Human NAD+
16	13.4	67.0	25	6	AX488345	AX488345 Sequence
17	13.4	67.0	26	6	AX117947	AX117947 Sequence
18	13.2	66.0	30	6	AX023694	AX023694 Sequence
19	13.2	66.0	31	6	AR105818	AR105818 Sequence
20	13.2	66.0	35	6	AR049511	AR049511 Sequence
21	13.2	66.0	36	6	AR003843	AR003843 Sequence
22	13.2	66.0	65	6	AX485913	AX485913 Sequence
23	13.2	66.0	65	6	E02972	E02972 DNA encodin
24	13.2	66.0	94	6	AR140872	AR140872 Sequence
25	13.2	66.0	94	6	AR150822	AR150822 Sequence
26	13.2	66.0	94	6	I65700	I65700 Sequence 60
27	13.2	66.0	94	6	I67932	I67932 Sequence 60
28	13.2	66.0	94	6	I90153	I90153 Sequence 60
29	12.8	64.0	20	6	AR203176	AR203176 Sequence
30	12.8	64.0	24	6	AR034080	AR034080 Sequence
31	12.8	64.0	25	6	E04736	E04736 PCR primer
32	12.8	64.0	27	6	AR099080	AR099080 Sequence
33	12.8	64.0	27	6	AR125428	AR125428 Sequence
34	12.8	64.0	27	6	AR127142	AR127142 Sequence
35	12.8	64.0	27	6	AR144700	AR144700 Sequence
36	12.8	64.0	27	6	AR151618	AR151618 Sequence
37	12.8	64.0	29	6	AR177299	AR177299 Sequence
38	12.8	64.0	29	6	BD003159	BD003159 Secretary
39	12.8	64.0	29	6	BD003183	BD003183 Secretary
40	12.8	64.0	29	6	I80078	I80078 Sequence 56
41	12.8	64.0	33	6	AR174978	AR174978 Sequence
42	12.8	64.0	33	6	AR208534	AR208534 Sequence
43	12.8	64.0	36	6	A38898	A38898 Sequence 6
44	12.8	64.0	36	6	A38899	A38899 Sequence 7
45	12.8	64.0	51	6	AX204531	AX204531 Sequence

ALIGNMENTS

RESULT 1	A89801	20 bp	DNA	linear	PAT 22-JAN-2000
LOCUS	Sequence 23 from Patent WO9832462.				
DEFINITION	A89801				
ACCESSION	A89801.1	GI:6738315			
VERSION	unidentified.				
KEYWORDS	unidentified.				
SOURCE	unclassified.				
ORGANISM	1 (bases 1 to 20)				
REFERENCE	Lipford,G.B. and Heeg,K.				
AUTHORS	PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND				
TITLE	OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION				
JOURNAL	Patent: WO 9832462-A 23 30-JUL-1998;				

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FEATURES
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    LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
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      /db_xref="taxon:32644"
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  Best Local Similarity 100.0%; Pred. No. 1.2e+02;
  Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AAGCGAAATGAATTGACT 20
    |||||||
Db 1 AAGCGAAATGAATTGACT 20

RESULT 2
LOCUS      A90888      20 bp      DNA      linear      PAT 22-JAN-2000
DEFINITION Sequence 23 from Patent EP0855184.
ACCESSION A90888
VERSION A90888.1 GI:6739338
KEYWORDS .
SOURCE .
  ORGANISM
    unidentified.
    unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Heeg,K.P. and Lipford,G.B.
TITLE Pharmaceutical composition comprising a polynucleotide and an
  antigen especially for vaccination
JOURNAL Patent: EP 0855184-A 23 29-JUL-1998;
  HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
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Db 1 AAGCGAAATGAATTGACT 20

RESULT 3
LOCUS      AX455545      20 bp      DNA      linear      PAT 06-JUL-2002
DEFINITION Sequence 22 from Patent WO0222809.
ACCESSION AX455545
VERSION AX455545.1 GI:21714613
KEYWORDS .
SOURCE .
  ORGANISM
    synthetic construct.
    synthetic construct
    artificial sequences.
REFERENCE 1
AUTHORS Bauer,S., Lipford,G. and Wagner,H.
TITLE Process for high throughput screening of cp9g-based
  immuno-agonist/antagonist
JOURNAL Patent: WO 0222809-A 22 21-MAR-2002;
  Coley Pharmaceutical GmbH (DE)
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    /note="Synthetic oligonucleotide"
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  Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AAGCGAAATGAATTGACT 20
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Db 1 AAGCGAAATGAATTGACT 20

RESULT 4
LOCUS      AX406763      22 bp      DNA      linear      PAT 14-JUN-2002
DEFINITION Sequence 19 from Patent WO0229044.
ACCESSION AX406763
VERSION AX406763.1 GI:21439688
KEYWORDS .
SOURCE .
  ORGANISM
    synthetic construct.
    synthetic construct
    artificial sequences.
REFERENCE 1
AUTHORS Hecker,M. and Wagner,A.H.
TITLE Modulation of the transcription of pro-inflammatory gene products
JOURNAL Patent: WO 0229044-A 19 11-APR-2002;
  Location/Qualifiers
  1..22
  /organism="synthetic construct"
  /db_xref="taxon:32630"
  /note="oligonucleotide"
BASE COUNT      10 a      2 c      6 g      4 t
ORIGIN

  Query Match      100.0%; Score 20; DB 6; Length 22;
  Best Local Similarity 100.0%; Pred. No. 1.2e+02;
  Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AAGCGAAATGAATTGACT 20
    |||||||
Db 3 AAGCGAAATGAATTGACT 22

RESULT 5
LOCUS      AX406764/c      22 bp      DNA      linear      PAT 15-JUN-2002
DEFINITION Sequence 20 from Patent WO0229044.
ACCESSION AX406764
VERSION AX406764.1 GI:21439689
KEYWORDS .
SOURCE .
  ORGANISM
    synthetic construct.
    synthetic construct
    artificial sequences.
REFERENCE 1
AUTHORS Hecker,M. and Wagner,A.H.
TITLE Modulation of the transcription of pro-inflammatory gene products
JOURNAL Patent: WO 0229044-A 20 11-APR-2002;
  Location/Qualifiers
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  /note="oligonucleotide"
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ORIGIN

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  Best Local Similarity 100.0%; Pred. No. 1.2e+02;
  Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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    |||||||
Db 20 AAGCGAAATGAATTGACT 1

RESULT 6
AX482985/c

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20
/note="single nucleotide polymorphism
Accession number cg43119818"

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	LOCUS	LOCUS		linear	PRI 30-OCT-1994
Db	22 AACAAAAATGAATTTCAGT 3				
Db	22 AACAAAAATGAATTTCAGT 3				

DEFINITION Human NAD+ ADP-ribosyltransferase (ADPRT) gene, exon 11.
ACCESSION M29774 M2953
VERSION M29774.1 GI:178176
KEYWORDS ADP-D-ribosyltransferase; NAD+ ADP-ribosyltransferase.
SEGMENT 11 of 23
SOURCE Human HeLa cell line, cDNA to mRNA, clones Hz[19,20,21,22,23,24].
and liver DNA.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 20; 70 to 89)
AUTHORS Auer,B., Nagl,U., Herzog,H., Schneider,R. and Schweiger,M.
TITLE Human nuclear NAD+ ADP-ribosyltransferase (polymerizing):
organization of the gene
JOURNAL DNA 8 (8), 575-580 (1989)
MEDLINE 90091744
PUBMED 2513174
REFERENCE 2 (bases 11 to 79)
AUTHORS Herzog,H., Zabel,B.U., Schneider,R., Auer,B., Hirsch-Kauffmann,M.
and Schweiger,M.
TITLE Human nuclear NAD+ ADP-ribosyltransferase: localization of the gene
on chromosome 1q41-q42 and expression of an active human enzyme in
Escherichia coli
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 86 (10), 3514-3518 (1989)
MEDLINE 89284454
PUBMED 2498872
COMMENT Draft entry and computer-readable sequence for [2] kindly submitted
by H.Herzog, 07-MAR-1989.
FEATURES
Location/Qualifiers
source 1..89
/organism="Homo sapiens"
/db_xref="taxon:9606"
/map="1q41-q42"
intron <1..10
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exon 11..79
/gene="PPOL"
/note="NAD+ ADP-ribosyltransferase, (EC 2.4.2.30; 5' end
put.); G00-119-508; putative"
/number=11
intron 80..>89
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BASE COUNT 33 a 17 c 19 g 20 t
ORIGIN About 0.6 kb after segment 10; chromosome 1q41-q42.
Query Match 68.0%; Score 13.6; DB 9; Length 89;
Best Local Similarity 80.0%; Pred. No. 8.le+04;
Matches 16; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
QY 1 AAGCGAAATGAATTGACT 20
II II IIIIIIIII III
Db 26 AAAAGAGATGAATTAAC 45

Search completed: December 12, 2002, 02:56:19
Job time : 323.116 secs

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GenCore version 5.1.3
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:04:50 ; Search time 92.087 Seconds
(without alignments)
440.192 Million cell updates/sec

Title: US-09-355-254F-9

Perfect score: 18
Sequence: 1 ggaatgacgttcctcgtg 18

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 2390332

Minimum DB seq length: 0
Maximum DB seq length: 100

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N_Geneseq_101002:*

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- 24: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	18	100.0	18	AAV46001	Immune adjuvant IL
2	18	100.0	18	AAV46001	Immune adjuvant IL
3	14.4	80.0	65	AAV46001	Murine Toll-like r
C 4	13.8	76.7	38	AAV46001	Rat spliced transc
C 5	13.8	76.7	89	AAV46001	Proteoglycan core
C 6	13.8	76.7	91	AAV46001	Bacillus clausii g
C 7	13.4	74.4	60	AAV46001	Human secreted pro
C 8	13.2	73.3	65	AAV46001	Human spliced tran
C 9	13.2	73.3	65	AAV46001	Rat spliced transc
					Mouse spliced tran

10	13.2	73.3	65	24	ABN58058
C 11	13.2	73.3	90	22	AAK22893
C 12	13.2	73.3	90	22	AAK22893
C 13	13.2	73.3	90	22	AAK49067
C 14	13.2	73.3	90	22	AAK49067
C 15	13.2	73.3	90	22	AAK49067
C 16	13.2	73.3	90	22	AAK49067
C 17	13.2	73.3	90	22	AAK49067
C 18	13.2	73.3	90	22	AAK49067
C 19	13.2	73.3	90	22	AAK49067
C 20	13.2	73.3	90	22	AAK49067
C 21	13.2	73.3	90	22	AAK49067
C 22	13.2	73.3	90	22	AAK49067
C 23	13.2	73.3	90	22	AAK49067
C 24	13.2	73.3	90	22	AAK49067
C 25	13.2	73.3	90	22	AAK49067
C 26	13.2	73.3	90	22	AAK49067
C 27	13.2	73.3	90	22	AAK49067
C 28	13.2	73.3	90	22	AAK49067
C 29	13.2	73.3	90	22	AAK49067
C 30	13.2	73.3	90	22	AAK49067
C 31	13.2	73.3	90	22	AAK49067
C 32	13.2	73.3	90	22	AAK49067
C 33	13.2	73.3	90	22	AAK49067
C 34	13.2	73.3	90	22	AAK49067
C 35	13.2	73.3	90	22	AAK49067
C 36	13.2	73.3	90	22	AAK49067
C 37	13.2	73.3	90	22	AAK49067
C 38	13.2	73.3	90	22	AAK49067
C 39	13.2	73.3	90	22	AAK49067
C 40	13.2	73.3	90	22	AAK49067
C 41	13.2	73.3	90	22	AAK49067
C 42	13.2	73.3	90	22	AAK49067
C 43	13.2	73.3	90	22	AAK49067
C 44	13.2	73.3	90	22	AAK49067
C 45	13.2	73.3	90	22	AAK49067

ALIGNMENTS

RESULT 1	AAV46001	standard; DNA; 18 BP.
ID	AAV46001	standard; DNA; 18 BP.
XX	AAV46001	
AC	AAV46001	
XX	AAV46001	
DT	16-OCT-1998	(first entry)
XX	Immune adjuvant IL-13.	
DE	Immune system; adjuvant; vaccine; cancer; prophylactic; pathogenicity; modulator; tolerance; regulator; helper cell; antigen; immunoglobulin; Ig class; autoimmune response; T-cell; B-cell; tumour; ss.	
XX	Class Bacteria.	
OS	Class Bacteria.	
XX	EP855184-A1.	
PN	EP855184-A1.	
XX	29-JUL-1998.	
PD	29-JUL-1998.	
XX	23-JAN-1997;	97EP-0101019.
PF	23-JAN-1997;	97EP-0101019.
XX	23-JAN-1997;	97EP-0101019.
PR	(HEEG/) HEEG K.	
XX	(LIPE/) LIPFORD G B.	
PA	(WAGN/) WAGNER H.	
XX	Heeg K, Lipford GB, Wagner H;	
PI	WPI; 1998-389630/34.	
XX		

PT Antigenic composition comprises polynucleotide fragment and antigen
 PT - used as vaccine to treat or prevent e.g. cancer or pathogen
 PT infections and to modulate immune response e.g. tolerance break and
 PT regulation of TH1/TH2 cells
 PS Example 5; Page 8; 28pp; English.
 XX
 CC AAV45993-V46019 are fragments of bacterial polynucleotides which are
 CC used as immune adjuvants for inclusion into vaccines to treat cancer and
 CC for prophylaxis and/or treatment of conditions caused by pathogenic
 CC micro-organisms. The polynucleotide is used for modulation of an immune
 CC response and the modulation is selected from the group break of
 CC tolerance, regulation of TH1/TH2 helper cell responses, switch of Ig
 CC classes, treatment of autoimmune responses and induction of tolerances.
 CC DNA oligomers are used to enhance the reactivity of immune cells to
 CC viral, bacterial and parasitic antigens, to break tolerance in anergic T
 CC and B cells e.g. against tumour antigens, as adjuvants in vaccination
 CC against tumour defined antigens and immunostimulatory substances in an
 CC immune response against tumours and to suppress immune reactions of the
 CC innate and acquired immune system. The composition is inexpensive and
 CC stable and does not cause lethal shock, which happens with prior art
 CC bacterial sequences.
 XX
 SQ Sequence 18 BP; 3 A; 4 C; 6 G; 5 T; 0 other;

Query Match 100.0%; Score 18; DB 19; Length 18;
 Best Local Similarity 100.0%; Pred. No. 2.3;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18
 Db 1 GGAATGACGTTCCCTGTG 18

RESULT 2

AAL39185
 ID AAL39185 standard; DNA; 18 BP.

XX AAL39185;

DT 05-SEP-2002 (first entry)

XX Murine Toll-like receptor related CpG DNA SEQ ID NO 60.

XX Murine Toll-like receptor; TLR9; TLR7; TLR8; ISNA; ds.

XX Unidentified.

XX WO200222809-A2.

XX 21-MAR-2002.

PF 17-SEP-2001; 2001WO-US29229.

XX 15-SEP-2000; 2000US-233035P.

PR 23-JAN-2001; 2001US-263657P.

PR 17-MAY-2001; 2001US-291726P.

XX 22-JUN-2001; 2001US-300210P.

XX (COLE-) COLEY PHARM GMBH.

PA Bauer S, Lipford G, Wagner H;

DR WPI; 2002-393964/42.

XX New isolated murine Toll-like receptor (TLR)9, TLR7, TLR8 polypeptides,
 PT useful for identifying species specificity of immunostimulatory nucleic
 PT acid and identifying immunostimulatory nucleic acids

PS Disclosure; Page 76; 195pp; English.

XX The invention relates to isolated murine Toll-like receptors (TLR)9,
 CC TLR7 and TLR8 polypeptides. These polypeptides comprise fully defined

CC sequences of 1032, 1050 or 1032 amino acids as given in specification, or
 CC their fragments, where TLR9, TLR7 and TLR8 polypeptides or their
 CC fragments have an amino acid sequence which is identical to human TLR9,
 CC TLR7 or TLR8 polypeptides or their fragment except for at least one amino
 CC acid of a murine TLR polypeptide. The isolated nucleic acids of the
 CC invention are useful for inhibiting TLR9 signalling activity in a cell.
 CC TLR7, TLR8 and TLR9 polypeptides are useful for identifying nucleic acid
 CC molecules which interact with a TLR polypeptide or its fragment. The
 CC TLR7, TLR8 or TLR9 polypeptides are also useful for identifying ISNA. The
 CC TLR7, TLR8 and TLR9 polypeptides are also useful for comparing TLR9
 CC signalling activity of a test compound (that is not a nucleic acid, and
 CC is a polypeptide or a part of a combinatorial library of compounds) with
 CC an ISNA. The TLR7, TLR8 and TLR9 polypeptides are also useful for
 CC identifying species specificity of an ISNA. The isolated nucleic acids of
 CC the invention are useful as probes or primers. This polynucleotide
 CC sequence represents DNA relating to the isolated Toll-like receptors of
 CC the invention.

SQ Sequence 18 BP; 3 A; 4 C; 6 G; 5 T; 0 other;

Query Match 100.0%; Score 18; DB 24; Length 18;
 Best Local Similarity 100.0%; Pred. No. 2.3;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18

Db 1 GGAATGACGTTCCCTGTG 18

RESULT 3

ABN30296

ID ABN30296 standard; DNA; 65 BP.

XX ABN30296;

DT 15-JUL-2002 (first entry)

XX Rat spliced transcript detection oligonucleotide SEQ ID NO:3044.

KW Human; mouse; rat; splice transcript; detection; RNA transcript;
 KW splice variant; transcriptome; oligonucleotide library; ss.

XX Rattus norvegicus.

XX WO200210449-A2.

XX 07-FEB-2002.

XX 20-JUL-2001; 2001WO-IB01903.

XX 28-JUL-2000; 2000US-221607P.

PR 02-MAY-2001; 2001US-287724P.

XX (COMP-) COMPUGEN INC.

XX Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;

XX WPI; 2002-257383/30.

XX New oligonucleotide libraries comprising oligonucleotides which
 PT selectively hybridize to mRNAs transcribed from a transcription unit of
 PT a genome, useful for detecting tissue-, pathology-, and
 PT developmental-specific genes

PS Example 1; SEQ ID 3044; 47pp; English.

XX The present invention describes oligonucleotide libraries for detecting
 CC messenger RNAs that populate a (sub-)transcriptome, where the
 CC (sub-)transcriptome comprises messenger RNAs transcribed from multiple
 CC transcription units that populate a genome. The library comprises
 CC several oligonucleotides, each capable of hybridising selectively to a
 CC set of messenger RNAs transcribed from a given transcription unit of
 CC the genome, which encodes one or more messenger RNA splice variants.

CC The oligonucleotide libraries are useful for detecting mRNAs from a
 CC biological sample, in expression profiling studies, in qualitatively or
 CC quantitatively characterising the corresponding transcriptome, and in
 CC detecting RNA transcripts and splice variants of human or animal
 CC transcripts. The libraries may also be used as specialised mini
 CC libraries to detect transcripts of a sub-transcriptome under a
 CC particular biological or pathological state, and so allowing the
 CC detection of tissue- and pathology-specific genes such as those genes
 CC only expressed in specific tissue under a specific pathological
 CC condition; to detect developmental specific genes; and to detect RNA
 CC transcripts and splice variants of a transcriptome of a patient suffering
 CC from a particular disorder. ABN27253 to ABN59589 represent
 CC oligonucleotide sequences from rats, humans and mice, which are used in
 CC the exemplification of the present invention.
 CC N.B. The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

SQ Sequence 65 BP; 18 A; 13 C; 16 G; 18 T; 0 other;

Query Match 80.0%; Score 14.4; DB 24; Length 65;
 Best Local Similarity 93.8%; Pred. No. 2.5e+02;
 Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 AATGACGTTCCCTGTG 18
 |||||
 Db 9 AATGACGTTCCCTGCG 24

RESULT 4

AAQ57712/c
 ID AAQ57712 standard; DNA; 38 BP.

AC AAQ57712;

DT 11-AUG-1994 (first entry)

DE Proteoglycan core glycoprotein antisense primer.

XX Neurocan; cell adhesion; leukocyte-endothelial cell recognition; lipid;
 KW tissue-related inflammation allergy; cellular; humoral; carbohydrate;
 KW hypersensitivity; trauma; neuronal development; cell transport; enzyme;
 KW infection; diagnosis; lectin; versican; aggrecan; gelsolin; saccharide;
 KW receptor; cell recognition; membrane cytoplasmic protein; nucleoside;
 KW ion; ss.

XX Synthetic.

XX Key Location/Qualifiers
 PH modified_base 28
 FT /*tag= a
 FT /label= Inosine

XX W09403601-A.

XX 17-FEB-1994.

XX 03-AUG-1993; 93WO-US07306.

XX 03-AUG-1992; 92US-0922911.

XX (UYNV) UNIV NEW YORK STATE.

XX Margolis RK, Margolis RU, Rauch U;

XX WPI; 1994-065690/08.

XX Eukaryotic neurocan polypeptide(s) with epidermal growth factor,
 PT lectin or complement binding activity - used in the diagnosis,
 PT treatment or research of hypersensitivity and allergic diseases
 XX Example 1; Fig 1B; 105pp; English.

XX

CC The sequences given in AAQ57711-12 represent primers which were used
 CC for the isolation of the neurocan cDNA. The neurocan protein
 CC has several biological activities, including cell adhesion, leukocyte-
 CC endothelial cell recognition, tissue-related inflammation allergies,
 CC cellular and/or humoral hypersensitivity, trauma, neuronal
 CC development, and cell transport and/or infection. Compositions
 CC containing them can be used as modulators of these conditions, and
 CC may be used as therapeutic, diagnostic, and/or research tools.
 CC Neurocan peptides can be used to mimic proteins, such as lectins,
 CC cell adhesion molecules, versicans, aggrecans or gelsolins, as
 CC receptor or effector subtypes. The protein can be used to treat
 CC diseases involving a qualitative or quantitative pathological
 CC abnormality of cell adhesion or leukocyte-endothelial cell recognition,
 CC or a functionally associated molecule such as a membrane cytoplasmic
 CC protein, lipid, carbohydrate, saccharide, nucleoside, enzyme or ion.
 XX

SQ Sequence 38 BP; 7 A; 9 C; 5 G; 10 T; 7 other;

Query Match 76.7%; Score 13.8; DB 15; Length 38;
 Best Local Similarity 70.6%; Pred. No. 5.2e+02;
 Matches 12; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGT 17

Db 38 GGAATGACGTTCCCTGT 22
 |||||

RESULT 5

ABK79561/c

ID ABK79561 standard; DNA; 89 BP.

AC ABK79561;

XX 13-AUG-2002 (first entry)

XX Bacillus clausii genomic sequence tag (GST) #2404.

XX Differential gene expression; genomic sequenced tag; GST;
 KW altered culture condition; environmental stress;
 KW physiological provocation; ds.

XX Bacillus clausii.

XX W0200229113-A2.

XX 11-APR-2002.

XX 05-OCT-2001; 2001WO-US31437.

XX 06-OCT-2000; 2000US-0680598.

XX 27-MAR-2001; 2001US-279526P.

XX (NOVO) NOVOZYMES BIOTECH INC.

XX (NOVO) NOVOZYMES AS.

XX Berka R, Clausen IG;

XX WPI; 2002-416684/44.

XX Monitoring differential expression of several genes in first Bacillus
 PT cell relative to expression of same genes in one or more second
 PT Bacillus cells, by using substrate containing Bacillus genomic
 PT sequenced tag array

XX Claim 11; SEQ ID NO 6852; 200pp; English.

XX The invention describes a method of monitoring differential expression of
 CC genes in a first Bacillus cell relative to expression of the genes in
 CC other Bacillus cells, comprising hybridising labelled nucleic acid probes
 CC isolated from Bacillus cells to a substrate containing array of Bacillus
 CC genomic sequenced tags (GST), examining the array, and determining
 CC relative gene expression by an observed hybridisation reporter signal of
 CC a spot in the array. The method is useful for measuring the expression of

CC the exemplification of the present invention.
CC N.B. The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 60 BP; 18 A; 5 C; 20 G; 17 T; 0 other;

Query Match 74.4%; Score 13.4; DB 24; Length 60;
Best Local Similarity 93.3%; Pred. No. 9e+02;
Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 AATGACGTTCCCTGT 17
||| ||||| ||||| |||||
Db 38 AATGACGTTCCCTGT 24

RESULT 8
ABN29564/C
ID ABN29564 standard; DNA; 65 BP.
XX
AC ABN29564;
XX
DT 15-JUL-2002 (first entry)
XX
DE Rat spliced transcript detection oligonucleotide SEQ ID NO:2312.
XX
KW Human; mouse; rat; splice transcript; detection; RNA transcript;
KW splice variant; transcriptome; oligonucleotide library; ss.
XX
OS Rattus norvegicus.
XX
PN WO200210449-A2.
XX
PD 07-FEB-2002.
XX
PF 20-JUL-2001; 2001WO-IB01903.
XX
PR 28-JUL-2000; 2000US-221607P.
PR 02-MAY-2001; 2001US-287724P.
XX
PA (COMP-) COMPUGEN INC.
XX
PI Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;
XX
DR WPI; 2002-257383/30.
XX
PT New oligonucleotide libraries comprising oligonucleotides which
PT selectively hybridize to mRNAs transcribed from a transcription unit of
PT a genome, useful for detecting tissue-, pathology-, and
PT developmental-specific genes -
XX
PS Example 1; SEQ ID 2312; 47pp; English.
XX
CC The present invention describes oligonucleotide libraries for detecting
CC messenger RNAs that populate a (sub-)transcriptome, where the
CC (sub-)transcriptome comprises messenger RNAs transcribed from multiple
CC transcription units that populate a genome. The library comprises
CC several oligonucleotides, each capable of hybridising selectively to a
CC set of messenger RNAs transcribed from a given transcription unit of
CC the genome, which encodes one or more messenger RNA splice variants.
CC The oligonucleotide libraries are useful for detecting mRNAs from a
CC biological sample, in expression profiling studies, in qualitatively or
CC quantitatively characterising the corresponding transcriptome, and in
CC detecting RNA transcripts and splice variants of human or animal
CC transcriptomes. The libraries may also be used as specialised mini
CC libraries to detect transcripts of a sub-transcriptome under a
CC particular biological or pathological state, and so allowing the
CC detection of tissue- and pathology-specific genes such as those genes
CC only expressed in specific tissue under a specific pathological
CC condition; to detect developmental specific genes; and to detect RNA
CC transcripts and splice variants of a transcriptome of a patient suffering
CC from a particular disorder. ABN27253 to ABN59589 represent
CC oligonucleotide sequences from rats, humans and mice, which are used in

CC the exemplification of the present invention.
CC N.B. The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 65 BP; 20 A; 17 C; 16 G; 12 T; 0 other;

Query Match 73.3%; Score 13.2; DB 24; Length 65;
Best Local Similarity 83.3%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18
||||| | ||||| |||||
Db 29 GGAATGTCCTTCCATGTG 12

RESULT 9
ABN58024
ID ABN58024 standard; DNA; 65 BP.
XX
AC ABN58024;
XX
DT 15-JUL-2002 (first entry)
XX
DE Mouse spliced transcript detection oligonucleotide SEQ ID NO:30772.
XX
KW Human; mouse; rat; splice transcript; detection; RNA transcript;
KW splice variant; transcriptome; oligonucleotide library; ss.
XX
OS Mus musculus.
XX
PN WO200210449-A2.
XX
PD 07-FEB-2002.
XX
PF 20-JUL-2001; 2001WO-IB01903.
XX
PR 28-JUL-2000; 2000US-221607P.
PR 02-MAY-2001; 2001US-287724P.
XX
PA (COMP-) COMPUGEN INC.
XX
PI Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;
XX
DR WPI; 2002-257383/30.
XX
PT New oligonucleotide libraries comprising oligonucleotides which
PT selectively hybridize to mRNAs transcribed from a transcription unit of
PT a genome, useful for detecting tissue-, pathology-, and
PT developmental-specific genes -
XX
PS Example 1; SEQ ID 30772; 47pp; English.
XX
CC The present invention describes oligonucleotide libraries for detecting
CC messenger RNAs that populate a (sub-)transcriptome, where the
CC (sub-)transcriptome comprises messenger RNAs transcribed from multiple
CC transcription units that populate a genome. The library comprises
CC several oligonucleotides, each capable of hybridising selectively to a
CC set of messenger RNAs transcribed from a given transcription unit of
CC the genome, which encodes one or more messenger RNA splice variants.
CC The oligonucleotide libraries are useful for detecting mRNAs from a
CC biological sample, in expression profiling studies, in qualitatively or
CC quantitatively characterising the corresponding transcriptome, and in
CC detecting RNA transcripts and splice variants of human or animal
CC transcriptomes. The libraries may also be used as specialised mini
CC libraries to detect transcripts of a sub-transcriptome under a
CC particular biological or pathological state, and so allowing the
CC detection of tissue- and pathology-specific genes such as those genes
CC only expressed in specific tissue under a specific pathological
CC condition; to detect developmental specific genes; and to detect RNA
CC transcripts and splice variants of a transcriptome of a patient suffering
CC from a particular disorder. ABN27253 to ABN59589 represent
CC oligonucleotide sequences from rats, humans and mice, which are used in

CC the exemplification of the present invention.
CC N.B. The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 65 BP; 15 A; 20 C; 19 G; 11 T; 0 other;

Query Match 73.3%; Score 13.2; DB 24; Length 65;
Best Local Similarity 83.3%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18
||||| |||||
Db 43 GGAAGGACGACCCCTGTG 60

RESULT 10
ABN58058
ID ABN58058 standard; DNA; 65 BP.
XX
AC ABN58058;
XX
DT 15-JUL-2002 (first entry)
XX
DE Mouse spliced transcript detection oligonucleotide SEQ ID NO:30806.

XX Human; mouse; rat; splice transcript; detection; RNA transcript;
KW splice variant; transcriptome; oligonucleotide library; ss...

XX Mus musculus.
XX WO200210449-A2.
XX
XX 07-FEB-2002.
XX
XX 20-JUL-2001; 2001WO-IB01903.
XX
XX 28-JUL-2000; 2000US-221607P.
XX
XX 02-MAY-2001; 2001US-287724P.
XX
XX (COMP-) COMPUEN INC.

XX Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;

XX WPI; 2002-257383/30.

XX New oligonucleotide libraries comprising oligonucleotides which
PT selectively hybridize to mRNAs transcribed from a transcription unit of
PT a genome, useful for detecting tissue-, pathology-, and
PT developmental-specific genes

XX Example 1; SEQ ID 30806; 47pp; English.

XX The present invention describes oligonucleotide libraries for detecting
CC messenger RNAs that populate a (sub-)transcriptome, where the
CC (sub-)transcriptome comprises messenger RNAs transcribed from multiple
CC transcription units that populate a genome. The library comprises
CC several oligonucleotides, each capable of hybridising selectively to a
CC set of messenger RNAs transcribed from a given transcription unit of
CC the genome, which encodes one or more messenger RNA splice variants.
CC The oligonucleotide libraries are useful for detecting mRNAs from a
CC biological sample, in expression profiling studies, in qualitatively or
CC quantitatively characterising the corresponding transcriptome, and in
CC detecting RNA transcripts and splice variants of human or animal
CC transcriptomes. The libraries may also be used as specialised mini
CC libraries to detect transcripts of a sub-transcriptome under a
CC particular biological or pathological state, and so allowing the
CC detection of tissue- and pathology-specific genes such as those genes
CC only expressed in specific tissue under a specific pathological
CC condition; to detect developmental specific genes; and to detect RNA
CC transcripts and splice variants of a transcriptome of a patient suffering
CC from a particular disorder. ABN27453 to ABN59589 represent
CC oligonucleotide sequences from rats, humans and mice, which are used in

CC the exemplification of the present invention.
CC N.B. The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 65 BP; 20 A; 17 C; 15 G; 13 T; 0 other;

Query Match 73.3%; Score 13.2; DB 24; Length 65;
Best Local Similarity 83.3%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18
||||| |||||
Db 15 GGAATGAAGTTCTCTGTG 32

RESULT 11
AAK22893/c
ID AAK22893 standard; DNA; 90 BP.
XX
AC AAK22893;
XX
DT 05-NOV-2001 (first entry)
XX
DE Human brain expressed single exon probe SEQ ID NO: 22884.

XX Human; brain expressed exon; gene expression analysis; probe;
KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;
KW epilepsy; cancer; ss.

XX Homo sapiens.

XX WO200157275-A2.

XX 09-AUG-2001.

XX 30-JAN-2001; 2001WO-US00667.

XX 04-FEB-2000; 2000US-0180312.

XX 26-MAY-2000; 2000US-0207456.

XX 30-JUN-2000; 2000US-0608408.

XX 03-AUG-2000; 2000US-0632366.

XX 21-SEP-2000; 2000US-0234687.

XX 27-SEP-2000; 2000US-0236359.

XX 04-OCT-2000; 2000GB-0024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.

XX Penn SG, Hanzel DK, Chen W, Rank DR;

XX WPI; 2001-483446/52.

XX Single exon nucleic acid probes for analyzing gene expression in human
PT brains

XX Example 4; SEQ ID NO: 22884; 650pp + Sequence Listing; English.

XX The present invention provides a number of single exon nucleic acid
CC probes which are derived from genomic sequences expressed in the human
CC brain. They can be used to measure gene expression in brain cell samples,
CC which may enable the diagnosis and improved treatment of nervous system
CC diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,
CC epilepsy and cancers. The present sequence is one of the probes of the
CC invention.

XX Sequence 90 BP; 18 A; 33 C; 15 G; 24 T; 0 other;

Query Match 73.3%; Score 13.2; DB 22; Length 90;
Best Local Similarity 83.3%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18
||||| |||||

```
Db      68 GGAAGAAGCTCCCTGTG 51
RESULT 12
AAK49067/c
ID      AAK49067 standard; DNA; 90 BP.
XX
XX      AAK49067;
AC
XX
XX      06-NOV-2001 (first entry)
DT
XX
XX      Human bone marrow expressed single exon probe SEQ ID NO: 23624.
DE
XX
XX      Human; bone marrow expressed exon; gene expression analysis; probe;
KW      microarray; cancer; leukaemia; lymphoma; myeloma; ss.
XX
XX      Homo sapiens.
OS
XX
XX      (MOLE-) MOLECULAR DYNAMICS INC.
PA
XX
XX      Penn SG, Hanzel DK, Chen W, Rank DR;
PI
XX      WPI; 2001-488900/53.
DR
XX
XX      Human genome-derived single exon nucleic acid probes useful for
PT      analyzing gene expression in human bone marrow -
XX
XX      Example 4; SEQ ID NO: 23624; 658pp + Sequence Listing; English.
PS
XX
XX      The present invention provides a number of single exon nucleic acid
CC      probes which are derived from genomic sequences expressed in the human
CC      bone marrow. They can be used to measure gene expression in bone marrow
CC      samples, which may enable the improved diagnosis and treatment of cancers
CC      such as lymphoma, leukaemia and myeloma. The present sequence is one of
CC      the probes of the invention.
XX
XX      Sequence 90 BP; 18 A; 33 C; 15 G; 24 T; 0 other;
SQ
Query Match      73.3%; Score 13.2; DB 22; Length 90;
Best Local Similarity 83.3%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY      1 GGAATGACGTTCCCTGTG 18
      ||||| | | |||||
Db      68 GGAAGAAGCTCCCTGTG 51

RESULT 13
AAI54894/c
ID      AAI54894 standard; DNA; 90 BP.
XX
XX      AAI54894;
AC
XX
XX      17-OCT-2001 (first entry)
DT
XX
XX      Probe #23580 used to measure gene expression in human placenta sample.
DE
XX
XX      Probe; microarray; human; placenta; antenatal diagnosis;
KW      genetic disorder; ss.

XX
XX      Homo sapiens.
OS
XX
XX      WO200157272-A2.
PN
XX
XX      09-AUG-2001.
PD
XX
XX      30-JAN-2001; 2001WO-US006563.
PF
XX
XX      04-FEB-2000; 2000US-0180312.
PR
XX      26-MAY-2000; 2000US-0207456.
PR
XX      30-JUN-2000; 2000US-0608408.
PR
XX      03-AUG-2000; 2000US-0632366.
PR
XX      21-SEP-2000; 2000US-0234687.
PR
XX      27-SEP-2000; 2000US-0236359.
PR
XX      04-OCT-2000; 2000GB-0024263.
XX
XX      (MOLE-) MOLECULAR DYNAMICS INC.
PA
XX
XX      Penn SG, Hanzel DK, Chen W, Rank DR;
PI
XX      WPI; 2001-48897/53.
DR
XX
XX      Human genome-derived single exon nucleic acid probes useful for
PT      analyzing gene expression in human placenta -
XX
XX      Claim 25; SEQ ID No 23580; 654pp; English.
PS
XX
XX      The present invention relates to single exon nucleic acid probes (SENP).
CC      The present sequence is one such probe. The probes are useful for
CC      producing a microarray for predicting, measuring and displaying gene
CC      expression in samples derived from human placenta. The probes are useful
CC      for antenatal diagnosis of human genetic disorders.
XX
XX      Sequence 90 BP; 18 A; 33 C; 15 G; 24 T; 0 other;
SQ
Query Match      73.3%; Score 13.2; DB 22; Length 90;
Best Local Similarity 83.3%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY      1 GGAATGACGTTCCCTGTG 18
      ||||| | | |||||
Db      68 GGAAGAAGCTCCCTGTG 51

RESULT 14
AAV23036/c
ID      AAV23036 standard; DNA; 28 BP.
XX
XX      AAV23036;
AC
XX
XX      30-JUL-1998 (first entry)
DT
XX
XX      HG3305S-28 primer used to amplify Hepatitis virus g gene sequences.
DE
XX
XX      Hepatitis g virus gene; diagnosis; treatment; Hepatitis g virus disease;
KW      PCR primer; ss.
XX
XX      Synthetic.
OS
XX      Hepatitis g virus.
XX
XX      JP10108685-A.
PN
XX
XX      28-APR-1998.
PD
XX
XX      10-AUG-1997; 97JP-0227387.
PF
XX
XX      10-AUG-1996; 96JP-0227639.
PR
XX
XX      (BMLB-) BML KK.
PA
XX
XX      WPI; 1998-304974/27.
XX
```

PT New hepatitis G virus gene - useful for diagnosing and treating
PT diseases caused by virus
XX
XX
XX Disclosure; Page 6; 128pp; Japanese.
XX
CC PCR primers AAV23018-74 were used to amplify and isolate new Hepatitis g
CC virus gene (see AAV23075-83 for gene fragments). RNA was synthesised
CC from the serum of nine patients judged positive for Hepatitis g virus
CC and cDNA synthesised from this RNA. The cDNA was used as a template in
CC several PCR reactions to isolate fragments of the new gene. The gene
CC may be useful for diagnosing and developing treatments for Hepatitis g
CC virus diseases.
XX
XX
SQ Sequence 28 BP; 8 A; 8 C; 5 G; 7 T; 0 other;
Query Match 72.2%; Score 13; DB 19; Length 28;
Best Local Similarity 100.0%; Pred. No. 1.4e+03; Length 28;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 4 ATGACGTTCCCTG 16
|||||
Db 28 ATGACGTTCCCTG 16
RESULT 15
AAAF74984
ID AAF74984 standard; DNA; 29 BP.
XX
XX AAF74984;
XX
XX 23-MAY-2001 (first entry)
DT
DE E. nidulans pPyrG sense primer pPyr3'.
XX
XX GlacA; promoter: pPyrG; PkIA; LacA; terminator; PCR primer; adapter;
KW protein expression; filamentous fungus; Escherichia coli;
KW Aspergillus niger; expression vector; ss.
XX
XX Emericella nidulans.
XX
XX WO200120007-A1.
PN
XX
XX 22-MAR-2001.
PD
XX
XX 13-SEP-2000; 2000WO-CA01084.
PF
XX
XX 13-SEP-1999; 99US-0153228.
PR
XX
XX (UYCO-) UNIV CONCORDIA.
PA
XX
XX Sillaots S, Martinez-Perez A, Tsang A, Storms R;
PI
XX
XX WPI; 2001-244813/25.
DR
XX
XX Expression vector for isolating filamentous fungi that expresses a
PT protein of interest at high levels, has selectable marker for fungi and
PT promoter operably linked to nucleic acid sequence encoding the protein
PT
XX
XX
XX Example 3; Page 33; 54pp; English.
XX
XX The present invention describes an expression vector (I) capable of
CC enabling a systematic analysis of gene expression and/or high-throughput
CC strain improvement screens in filamentous fungi, comprising a selectable
CC marker for the filamentous fungi and a promoter operably linked to a
CC sequence encoding a protein of interest or its part. Also described is a
CC filamentous fungi strain (II), enabling the expression of a protein of
CC interest to greater than 100, preferably greater than 1000 times
CC compared to the level in a parent strain. (I) is useful for isolating a
CC filamentous fungi, in particular Aspergillus niger that expresses a
CC protein of interest at high levels required for industrial-scale protein
CC production. (I) increases the expression of the protein of interest by
CC 100 times, preferably 1000 times, as compared to the level of expression

CC in a control filamentous fungi strain. (I) increases the efficiency and
CC reduces the time required for isolating strains that express proteins of
CC interest at high levels. AAF74980 to AAF74997 represent oligonucleotide
CC sequences used in an example from the present invention.
XX
XX
SQ Sequence 29 BP; 8 A; 8 C; 5 G; 8 T; 0 other;
Query Match 71.1%; Score 12.8; DB 22; Length 29;
Best Local Similarity 87.5%; Pred. No. 1.8e+03;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 2 GAATGACGTTCCCTGT 17
|||||
Db 2 GAATGACGTTCCCTTT 17
Search completed: December 12, 2002, 01:36:19
Job time : 96.087 secs

GenCore version 5.1.3
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:06:16 ; Search time 318.116 Seconds
(without alignments)
1829.698 Million cell updates/sec

Title: US-09-355-254F-23
Perfect score: 20
Sequence: 1 caggcataaacggttcogtag 20

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0
Maximum DB seq length: 100

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : GenEmbl:

1: gb_ba:*
2: gb_hgt:*
3: gb_in:*
4: gb_om:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vi:*
15: em_ba:*
16: em_fun:*
17: em_hum:*
18: em_in:*
19: em_mu:*
20: em_om:*
21: em_or:*
22: em_ov:*
23: em_pat:*
24: em_ph:*
25: em_pi:*
26: em_ro:*
27: em_sts:*
28: em_un:*
29: em_vi:*
30: em_htg_hum:*
31: em_htg_inv:*
32: em_htg_other:*
33: em_htg_mus:*
34: em_htg_pln:*
35: em_htg_rod:*
36: em_htg_mam:*
37: em_htg_vrt:*
38: em_sy:*
39: em_htgo_hum:*
40: em_htgo_mus:*
41: em_htgo_other:*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	20	100.0	20	6	A9802	A9802 Sequence 24
2	20	100.0	20	6	A90889	A90889 Sequence 24
3	20	100.0	20	6	AX455564	AX455564 Sequence
4	15.2	76.0	25	6	AX196810	AX196810 Sequence
5	13.6	68.0	24	6	AX445605	AX445605 Sequence
6	13.4	67.0	31	6	BD002472	BD002472 Gene comp
7	13	65.0	20	6	E07207	E07207 PCR primer
8	12.8	64.0	33	6	AX262268	AX262268 Sequence
9	12.8	64.0	35	6	AX262303	AX262303 Sequence
10	12.8	64.0	39	6	AX262285	AX262285 Sequence
11	12.8	64.0	65	6	AX486035	AX486035 Sequence
12	12.8	64.0	97	6	I44826	I44826 Sequence 1
13	12.6	63.0	33	6	A32217	A32217 Synthetic m
14	12.6	63.0	50	6	AX103395	AX103395 Sequence
15	12.2	61.0	30	6	AX374961	AX374961 Sequence
16	12.2	61.0	31	6	AR106260	AR106260 Sequence
17	12.2	61.0	31	6	E61282	E61282 Method for
18	12.2	61.0	37	6	AR145069	AR145069 Sequence
19	12.2	61.0	37	6	AR168079	AR168079 Sequence
20	12.2	61.0	37	6	AR169792	AR169792 Sequence
21	12.2	61.0	37	6	AR204850	AR204850 Sequence
22	12	60.0	33	6	A48801	A48801 Sequence 2
23	12	60.0	36	6	AX247485	AX247485 Sequence
24	12	60.0	39	6	AX052711	AX052711 Sequence
25	12	60.0	51	6	AX159959	AX159959 Sequence
26	12	60.0	63	6	AX482092	AX482092 Sequence
27	12	60.0	72	9	HSU91302	HSU91302 Homo sapien
28	12	60.0	90	9	HSC1CHX13	225758 Homo sapien
29	11.8	59.0	24	6	AX445387	AX445387 Sequence
30	11.8	59.0	26	6	AR125091	AR125091 Sequence
31	11.8	59.0	31	6	BD002473	BD002473 Gene comp
32	11.8	59.0	33	6	AR008899	AR008899 Sequence
33	11.8	59.0	33	6	AR087594	AR087594 Sequence
34	11.8	59.0	35	6	A84225	A84225 Sequence 4
35	11.8	59.0	38	6	AR038218	AR038218 Sequence
36	11.8	59.0	39	6	AX080158	AX080158 Sequence
37	11.8	59.0	61	6	AX080156	AX080156 Sequence
38	11.8	59.0	65	6	AX484924	AX484924 Sequence
39	11.8	59.0	100	14	S82444	S82444 [3' region,
40	11.8	59.0	100	14	S82445	S82445 [3' region,
41	11.8	59.0	100	14	S82446	S82446 [3' region,
42	11.6	58.0	24	6	AX443631	AX443631 Sequence
43	11.6	58.0	25	6	AX447613	AX447613 Sequence
44	11.6	58.0	28	6	AR160111	AR160111 Sequence
45	11.6	58.0	35	6	AR084174	AR084174 Sequence

ALIGNMENTS

RESULT 1
A98802
LOCUS A98802
DEFINITION Sequence 24 from Patent WO9832462.
ACCESSION A98802
VERSION A98802.1 GI:6738316
KEYWORDS
SOURCE unidentified.
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Lipford,G.B. and Heeg,K.
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
JOURNAL Patent: WO 9832462-A 24 30-JUL-1998;

LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
Location/Qualifiers

1..20
/organism="unidentified"
/db_xref="taxon:32644" 4 t

BASE COUNT 5 a 5 c 6 g 4 t

ORIGIN

Query Match 100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 9.7; Mismatches 0; Indels 0; Gaps 0;
Matches 20; Conservative 0;

QY 1 CAGGCATACGGTTCGGTAG 20

Db 1 CAGGCATACGGTTCGGTAG 20

RESULT 2

A90889

LOCUS

DEFINITION Sequence 24 from Patent EP0855184. 20 bp DNA linear PAT 22-JAN-2000

ACCESSION A90889

VERSION A90889.1 GI:6739343

KEYWORDS

SOURCE

ORGANISM

unidentified.

unclassified.

REFERENCE 1 (bases 1 to 20)

AUTHORS Heeg,K.P. and Lipford,G.B.

TITLE Pharmaceutical composition comprising a polynucleotide and an

JOURNAL antigen especially for vaccination

Patent: EP 0855184-A 24 29-JUL-1998;

HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)

FEATURES

Source

1..20

/organism="unidentified"

/db_xref="taxon:32644" 4 t

BASE COUNT 5 a 5 c 6 g 4 t

ORIGIN

Query Match 100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 9.7; Mismatches 0; Indels 0; Gaps 0;
Matches 20; Conservative 0;

QY 1 CAGGCATACGGTTCGGTAG 20

Db 1 CAGGCATACGGTTCGGTAG 20

RESULT 3

AX455564

LOCUS

DEFINITION Sequence 41 from Patent WO222809. 20 bp DNA linear PAT 06-JUL-2002

ACCESSION AX455564

VERSION AX455564.1 GI:21714632

KEYWORDS

SOURCE

ORGANISM

synthetic construct.

artificial sequences.

REFERENCE 1

AUTHORS Bauer, S., Lipford, G. and Wagner, H.

TITLE Process for high throughput screening of cpq-based

JOURNAL immuno-agonist/antagonist

Patent: WO 0222809-A 41 21-MAR-2002;

Coley Pharmaceutical GmbH (DE)

FEATURES

Source

1..20

/organism="synthetic construct"

/db_xref="taxon:32630"

/note="Synthetic oligonucleotide"

BASE COUNT 5 a 5 c 6 g 4 t

ORIGIN

Query Match 100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 9.7; Mismatches 0; Indels 0; Gaps 0;
Matches 20; Conservative 0;

QY 1 CAGGCATACGGTTCGGTAG 20

Db 1 CAGGCATACGGTTCGGTAG 20

RESULT 4

AX196810/c

LOCUS

DEFINITION Sequence 517 from Patent WO0151627. 25 bp DNA linear PAT 07-SEP-2001

ACCESSION AX196810

VERSION AX196810.1 GI:15387016

KEYWORDS

SOURCE

ORGANISM

soybean.

Glycine max

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;

Rosidae; eurosids I; Fabales; Fabaceae; Papilionoideae; Phaseoleae;

Glycine.

REFERENCE 1 (bases 1 to 25)

AUTHORS Haug, B.M., Wang, M.L., Parsons, J.D. and Parnell, L.D.

TITLE Nucleic acid molecules and other molecules associated with soybean

JOURNAL cyst nematode resistance

Patent: WO 0151627-A 517 19-JUL-2001;

MONSANTO COMPANY (US)

FEATURES

Source

1..25

/organism="Glycine max"

/db_xref="taxon:3847"

/note="Seq ID: 240017_region_G3_6395_12_Forward_Primer"

BASE COUNT 6 a 6 c 6 g 7 t

ORIGIN

Query Match 76.0%; Score 15.2; DB 6; Length 25;

Best Local Similarity 85.0%; Pred. No. 3.6e+03;

Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CAGGCATACGGTTCGGTAG 20

Db 22 CAGGCACCGGTTTCAGTAG 3

RESULT 5

AX445605

LOCUS

DEFINITION Sequence 2060 from Patent WO0216649. 24 bp DNA linear PAT 03-JUL-2002

ACCESSION AX445605

VERSION AX445605.1 GI:21692886

KEYWORDS

SOURCE

ORGANISM

synthetic construct.

artificial sequences.

REFERENCE 1

AUTHORS Gunderson, K.

TITLE Probes and decoder oligonucleotides

JOURNAL Patent: WO 0216649-A 2060 28-FEB-2002;

ILLUMINA, Inc. (US)

FEATURES

Source

1..24

/organism="synthetic construct"

/db_xref="taxon:32630"

/note="Computer Generated Probe Sequence."

BASE COUNT 6 a 5 c 9 g 4 t

ORIGIN

Query Match 68.0%; Score 13.6; DB 6; Length 24;

Best Local Similarity 80.0%; Pred. No. 2.6e+04;

Matches 16; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 CAGGCATACGGTTCGGTAG 20


```

Db      2  CAGGCATGACAGTTCGGTAG 21
|||||  ||  ||  |||||
RESULT 6
BD002472
LOCUS   31 bp      DNA      linear      PAT 31-JAN-2002
DEFINITION
Gene composition and method.
ACCESSION
BD002472
VERSION
BD002472.1 GI:18630433
KEYWORDS
JP 2000245487-A/138.
SOURCE
unidentified.
ORGANISM
unclassified.
REFERENCE
1 (bases 1 to 31)
AUTHORS
Sha,N., Walinton,J. and Patel,N.
TITLE
Gene composition and method
JOURNAL
Patent: JP 2000245487-A 138 12-SEP-2000;
AFIMETRICS INC
COMMENT
OS Unknown
PN JP 2000245487-A/138
PD 12-SEP-2000
PF 27-JAN-2000 JP 2000019392
PR 27-JAN-1999 US 09/238.402
PI NIRA SHA,JANET WALINTON,NIRA PATEL
PC C12N15/09,C12Q1/68,C12N15/00
CC
FH Key      Location/Qualifiers
FT source   1..31 /organism='Unknown'
FT
FEATURES
source
Location/Qualifiers
1..31
/organism="unidentified"
/db_xref="taxon:32644"
BASE COUNT  10 a  6 c  5 g  9 t  1 others
ORIGIN
1  CAGGCATGACAGTTCGG 17
|||||  ||  |||||
Query Match 67.0%; Score 13.4; DB 6; Length 31;
Best Local Similarity 82.4%; Pred. No. 3.3e+04;
Matches 14; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1  CAGGCATGACAGTTCGG 17
|||||  ||  |||||
Db      15  CRGGCAAACTGTTCCG 31

RESULT 7
E07207/c
LOCUS   20 bp      DNA      linear      PAT 29-SEP-1997
DEFINITION
PCR primer to detect Acholeplasma sp.
ACCESSION
E07207
VERSION
E07207.1 GI:2175348
KEYWORDS
JP 1994098800-A/2.
SOURCE
unidentified.
ORGANISM
unclassified.
REFERENCE
1 (bases 1 to 20)
AUTHORS
Nakagawa,T., Uemori,T., Asada,K., Katou,I. and Harasawa,A.
TITLE
METHOD FOR DETECTING ACHOLEPLASMA
JOURNAL
Patent: JP 1994098800-A 2 12-APR-1994;
TAKARA SHUZO CO LTD
COMMENT
OS None
OC Artificial sequences.
PN JP 1994098800-A/2
PD 12-APR-1994
PF 21-SEP-1992 JP 1992274830
PI NAKAGAWA TOMOKO, UEMORI TAKASHI, ASADA KIYOZOU, PI KATOU IKUNOSHIN,
PI HARASAWA AKIRA
PC C12Q1/68,C12N15/11,C12Q1/04,(C12Q1/04,C12R1:01); CC
strandedness: Single;
CC topology: Linear;
FH Key      Location/Qualifiers

```

```

FH      source   1..20 /organism='Artificial sequences'.
FT
FEATURES
source
Location/Qualifiers
1..20
/organism="unidentified"
/db_xref="taxon:32644"
BASE COUNT  4 a  7 c  3 g  4 t  2 others
ORIGIN
1  CAGGCATGACAGTTCGGTAG 20
|||||  ||  |||||
Query Match 65.0%; Score 13; DB 6; Length 20;
Best Local Similarity 76.5%; Pred. No. 5.4e+04;
Matches 13; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 4  GCATAACGGTTCGGTAG 20
|||||  ||  |||||
Db      18  GGATCAGGTTCSGTAR 2

RESULT 8
AX262268
LOCUS   33 bp      DNA      linear      PAT 26-OCT-2001
DEFINITION
Sequence 24 from Patent WO0173052.
ACCESSION
AX262268
VERSION
AX262268.1 GI:16511217
KEYWORDS
synthetic construct.
SOURCE
synthetic construct.
ORGANISM
artificial sequences.
REFERENCE
1
AUTHORS
Mchenry,C.S.
TITLE
Thermophilic polymerase III holoenzyme
JOURNAL
Patent: WO 0173052-A 24 04-OCT-2001;
Mchenry, Charles S. (US)
FEATURES
source
Location/Qualifiers
1..33
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="ATG forward primer P118-S85"
BASE COUNT  13 a  10 c  5 g  5 t
ORIGIN
1  CATAACGGTTCGGTAG 20
|||||  ||  |||||
Query Match 64.0%; Score 12.8; DB 6; Length 33;
Best Local Similarity 87.5%; Pred. No. 6.8e+04;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5  CATAACGGTTCGGTAG 20
|||||  ||  |||||
Db      11  CATAACGGTTCACAAG 26

RESULT 9
AX262303
LOCUS   35 bp      DNA      linear      PAT 26-OCT-2001
DEFINITION
Sequence 59 from Patent WO0173052.
ACCESSION
AX262303
VERSION
AX262303.1 GI:16511248
KEYWORDS
synthetic construct.
SOURCE
synthetic construct.
ORGANISM
artificial sequences.
REFERENCE
1
AUTHORS
Mchenry,C.S.
TITLE
Thermophilic polymerase III holoenzyme
JOURNAL
Patent: WO 0173052-A 59 04-OCT-2001;
Mchenry, Charles S. (US)
FEATURES
source
Location/Qualifiers
1..35
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="forward/sense ATG primer P118-S78cla2"
BASE COUNT  15 a  7 c  6 g  7 t
ORIGIN
1  CATAACGGTTCACAAG 26

```

Query Match	64.0%	Score 12.8;	DB 6;	Length 35;
Best Local Similarity	87.5%;	Pred. No. 6.8e+04;		
Matches 14;	Conservative	0;	Mismatches 2;	Indels 0;
Qy	5	CATAACGGTTCCTGAG	20	
Db	17	CATAACGGTTCCTGAG	32	

```
FEATURES
  source      Location/Qualifiers
1..50
/organism="unidentified adenovirus"
/db_xref="taxon:10535"
misc_feature 1..50
              /note="BAD"
BASE COUNT   15 a 10 c 10 g 15 t
ORIGIN

Query Match      63.0%; Score 12.6; DB 6; Length 50;
Best Local Similarity 78.9%; Pred. No. 8.7e+04;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 CAGGCATAACGGTTCGGTA 19
    |||||
Db 33 CAGCCAAAGTGTTCGGTA 15

RESULT 15
AX374961
LOCUS      AX374961          30 bp      DNA      linear      PAT 01-MAR-2002
DEFINITION Sequence 14 from Patent WO0210425.
ACCESSION  AX374961
VERSION    AX374961.1 GI:19169806
KEYWORDS   .
SOURCE     synthetic construct.
           synthetic construct
           artificial sequences.
REFERENCE  1
AUTHORS   Porto,D. and Sauer,M.
TITLE     Ascorbic acid production from yeasts
JOURNAL   Patent: WO 0210425-A 14 07-FEB-2002;
          Biopolo S.C.A.R.L. (IT)
FEATURES   Location/Qualifiers
1..30
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Forward PCR Primer for L-galactono-1,4-lactone
          dehydrogenase from A. thaliana"
BASE COUNT   8 a 9 c 6 g 7 t
ORIGIN

Query Match      61.0%; Score 12.2; DB 6; Length 30;
Best Local Similarity 82.4%; Pred. No. 1.4e+05;
Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 AGGCATAACGGTTCGGT 18
    |||||
Db 6 AGCCCTAAATGTTCGGT 22

Search completed: December 12, 2002, 02:56:25
Job time : 324.116 secs
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GenCore version 5.1.3
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OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:04:50 ; Search time 102.319 Seconds
(without alignments)
440.192 Million cell updates/sec

Title: US-09-355-254F-8

Perfect score: 20
Sequence: 1 gattcgctgacgtcagagag 20

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 2390332

Minimum DB seq length: 0
Maximum DB seq length: 100

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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- 2: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1981.DAT.*
- 3: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1982.DAT.*
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- 22: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
- 23: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
- 24: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20	100.0	20	AAV46000	Immune adjuvant AP
2	20	100.0	20	AAZ28189	Chlamydia trachoma
3	20	100.0	20	AAZ299172	Inflammatory cardi
4	20	100.0	20	AAZ39177	Murine Toll-like r
5	20	100.0	26	AAZ76047	CAMP response elem
6	20	100.0	27	AAT85832	CRE oligonucleotide
7	20	100.0	27	AAV82454	ATF comp oligonucle
8	20	100.0	27	AAV08336	CRE element coding
9	20	100.0	27	AAI70581	transcription fact

10	20	100.0	27	AAH77396	Cyclic AMP respons
11	20	100.0	27	AAH77397	Cyclic AMP respons
12	20	100.0	27	AAF87956	Cyclic AMP respons
13	20	100.0	27	AAF87957	Cyclic AMP respons
14	20	100.0	27	AAF76267	CAMP response elem
15	20	100.0	27	ABA92274	CRE binding site o
16	20	100.0	27	ABA05538	Cyclic-AMP respons
17	19	95.0	20	AAV45997	Immune adjuvant CR
18	16.4	82.0	28	ABK14052	Cyclic AMP respons
19	16.4	82.0	32	ABF77813	CREB probe derived
20	16.4	82.0	37	AAV04084	Somatostatin gene
21	16.4	82.0	50	AAQ69701	Human somatostatin
22	16.4	82.0	50	AAI64163	Human somatostatin
23	16.4	82.0	50	AAI17451	Test sequence from
24	16.4	82.0	50	ABK82942	DNA binding molecu
25	15.8	79.0	20	AAV45999	Immune adjuvant CR
26	14.8	74.0	95	AAC30099	Human secreted pro
27	14.4	72.0	33	ABK87820	Somatostatin promo
28	14.4	72.0	33	ABK87821	Somatostatin promo
29	14.4	72.0	83	AA598555	Human Pleckstrin h
30	14.2	71.0	41	AAV50622	Brassica sp. polym
31	14.2	71.0	41	ABL96051	Brassica polymorph
32	14.2	71.0	60	ABN49177	Human spliced tran
33	13.8	69.0	65	ABN57982	Mouse spliced tran
34	13.6	68.0	51	AAI78809	Human silent SNP c
35	13.4	67.0	96	AAI12454	Human secreted pro
36	13.2	66.0	22	AAV04086	PEPCK gene CAMP re
37	13.2	66.0	25	AAH39807	SNP specific SNPE
38	13.2	66.0	47	AAZ69295	Human map-related
39	13.2	66.0	51	AAH39808	Human SNP flanking
40	13.2	66.0	60	ABN45698	Human spliced tran
41	13.2	66.0	62	AAI66152	Hepatitis E virus
42	13.2	66.0	65	ABN56672	Mouse spliced tran
43	13.2	66.0	90	ABK36416	HIV DNA encoding N
44	13	65.0	26	AAV46019	Immune adjuvant CR
45	12.8	64.0	21	AAV41019	Primer PL2FRARA:12

ALIGNMENTS

RESULT 1
AAV46000
ID AAV46000 standard; DNA; 20 BP.

AC AAV46000;

DT 16-OCT-1998 (first entry)

DE Immune adjuvant AP-1 #1.

KW Immune system; adjuvant; vaccine; cancer; prophylactic; pathogenicity;
modulator; tolerance; regulator; helper cell; antigen; immunoglobulin;
KW Ig class; autoimmune response; T-cell; B-cell; tumour; ss.

OS Class Bacteria..

PN EP855184-A1.

XX 29-JUL-1998.

PF 23-JAN-1997; 97EP-0101019.

PR 23-JAN-1997; 97EP-0101019.

PA (HEG/) HEG K.

PA (LIPE/) LIPFORD G B.

PA (WAGN/) WAGNER H.

PI Heeg K, Lipford GB, Wagner H;

XX WPI; 1998-389630/34.

PT Antigenic composition comprises polynucleotide fragment and antigen
PT - used as vaccine to treat or prevent e.g. cancer or pathogen
PT infections and to modulate immune response e.g. tolerance break and
PT regulation of TH1/TH2 cells
XX
PS Example 3; Page 7; 28pp; English.
XX
CC AAV45993-V46019 are fragments of bacterial polynucleotides which are
CC used as immune adjuvants for inclusion into vaccines to treat cancer and
CC for prophylaxis and/or treatment of conditions caused by pathogenic
CC micro-organisms. The polynucleotide is used for modulation of an immune
CC response and the modulation is selected from the group break of
CC tolerance, regulation of TH1/TH2 helper cell responses, switch of Ig
CC classes, treatment of autoimmune responses and induction of tolerances.
CC DNA oligomers are used to enhance the reactivity of immune cells to
CC viral, bacterial and parasitic antigens, to break tolerance in anergic T
CC and B cells e.g. against tumour antigens, as adjuvants in vaccination
CC against tumour defined antigens and immunostimulatory substances in an
CC immune response against tumours and to suppress immune reactions of the
CC innate and acquired immune system. The composition is inexpensive and
CC stable and does not cause lethal shock, which happens with prior art
CC bacterial sequences.
XX
SQ Sequence 20 BP; 5 A; 4 C; 7 G; 4 T; 0 other;
Query Match 100.0%; Score 20; DB 19; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.48;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GATTGCTGACGTCAGAGAG 20
Db 1 GATTGCTGACGTCAGAGAG 20
IIIIIIIIIIIIIIIIIIII
RESULT 2
AAZ28189
ID AAZ28189 standard; DNA; 20 BP.
XX
AC AAZ28189;
XX
DT 20-DEC-1999 (first entry)
XX
DE Chlamydia trachomatis outer membrane protein gene-derived CpG oligo 2.
XX
KW Heart disease; inflammatory; autoimmune; cardiomyopathy; adjuvant;
KW CpG motif; vaccine; ds.
XX
OS Synthetic.
OS Chlamydia trachomatis.
XX
FH Key Location/Qualifiers
FH modified_base 1..20
FT /tag= a
FT /mod_base= OTHER
FT /note= "OTHER = phosphorothioate linkage"
XX
PN US5962636-A.
XX
PD 05-OCT-1999.
XX
PF 12-AUG-1998; 98US-0133774.
XX
PR 12-AUG-1998; 98US-0133774.
XX
PA (AMGE-) AMGEN CANADA INC.
XX
PI Bachmaier K, Hessel AJ, Penninger JM, Neu N;
XX WPI; 1999-589735/50.
XX
PT Peptides that induce or suppress inflammatory cardiomyopathy
XX Example 2 a-Column 25; 17pp; English.

XX This sequence represents DNA encoding Chlamydia trachomatis 60 kD outer
CC membrane protein (OMP) gene-derived CpG oligonucleotide 2. This
CC oligonucleotide contains a CpG motif. It was tested for its ability to
CC act as an adjuvant for the M7A-alpha peptide (AAV42723), which can
CC induce inflammatory cardiomyopathy (ICM) in mice. It was found to act as
CC a potent immunostimulant, whereas a oligonucleotide from the same
CC source which did not contain a CpG motif (AAZ28193) was hardly effective
CC as an adjuvant. Inflammatory cardiomyopathy peptides (AAV42723, AAV42725-
CC AAV42725-V42731) can be used with such an adjuvant and an excipient in a
CC vaccine for decreasing ICM.
XX
SQ Sequence 20 BP; 5 A; 4 C; 7 G; 4 T; 0 other;
Query Match 100.0%; Score 20; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.48;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GATTGCTGACGTCAGAGAG 20
Db 1 GATTGCTGACGTCAGAGAG 20
IIIIIIIIIIIIIIIIIIII
RESULT 3
AAZ99172
ID AAZ99172 standard; DNA; 20 BP.
XX
AC AAZ99172;
XX
DT 21-JUN-2000 (first entry)
XX
DE Inflammatory cardiomyopathy immunostimulatory oligonucleotide #1.
XX
KW Cardiant; murine alpha myosin heavy chain; inflammatory myocarditis;
KW autoimmune inflammatory cardiomyopathy; Chlamydia; antibody; vaccine;
KW hybridization probe; immunostimulatory; ss.
XX
OS Synthetic.
XX
PN US6034230-A.
XX
PD 07-MAR-2000.
XX
PF 03-MAY-1999; 99US-0303862.
XX
PR 12-AUG-1998; 98US-0133774.
XX
PA (AMGE-) AMGEN CANADA INC.
XX
PI Neu N, Penninger JM, Bachmaier K, Hessel AJ;
XX WPI; 2000-255712/22.
XX
PT DNA molecules encoding novel myocardial peptides used for inhibiting
PT and inducing inflammatory cardiomyopathy in vivo
XX
PS Disclosure; Column 17; 17pp; English.
XX
CC The invention relates to the isolation of sequences coding for peptide
CC sequences derived from bacteria and viruses which may cause inflammatory
CC cardiomyopathy. The peptide sequences are searched based on the sequence
CC of the M7A peptides derived from the murine alpha myosin heavy chain
CC polypeptide. The consensus sequence of the murine M7A-alpha/beta peptides
CC (Y83813) was used to search the PIR public database for similar bacterial
CC and viral sequences able to cause inflammatory cardiomyopathy. The screen
CC isolated the peptides Y83814-Y83819 and their corresponding coding
CC sequences 299164-299169. The peptides encoded by the DNAs are used, alone
CC or in conjunction with other therapeutics, for inducing or inhibiting
CC inflammatory cardiomyopathy in vivo, where the cardiomyopathy is
CC autoimmune inflammatory cardiomyopathy, and inflammatory cardiomyopathy
CC caused by Chlamydia or other bacterial or viral infections that cause
CC inflammatory cardiomyopathy. The oligonucleotides 299172-299176 were
CC shown to increase the immunogenicity of the immunostimulatory peptides

CC	when injected simultaneously. The peptides may also be used for
CC	increasing inflammatory myocarditis in a mammal. Antibodies against the
CC	peptides and the peptides themselves are used for measuring the risk of
CC	inflammatory cardiomyopathy in a mammal. The peptides may also be used
CC	in vaccines. Nucleic acids encoding the peptides may be used as
CC	hybridization probes, e.g. in diagnostic assays to test for the
CC	presence of Chlamydia DNA.
XX	
SQ	Sequence 20 BP; 5 A; 4 C; 7 G; 4 T; 0 other;
	Query Match 100.0%; Score 20; DB 21; Length 20;
	Best Local Similarity 100.0%; Pred. No. 0.48;
	Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	1 GATTGCCTGACGTCAGAGAG 20
DB	1 GATTGCCTGACGTCAGAGAG 20
RESULT 4	
AAL391177	ID AAL391177 standard; DNA; 20 BP.
XX	AC AAL391177;
XX	XX
DT	05-SEP-2002 (first entry)
XX	Murine Toll-like receptor related CpG DNA SEQ ID No 52.
DE	XX
XX	Murine Toll-like receptor; TLR9; TLR7; TLR8; ISNA; ds.
KW	Unidentified.
XX	OS
XX	WO200222809-A2.
PN	XX
XX	21-MAR-2002.
PD	XX
XX	17-SEP-2001; 2001WO-US29229.
PF	XX
PR	15-SEP-2000; 2000US-233035P.
PR	23-JAN-2001; 2001US-263657P.
PR	17-MAY-2001; 2001US-291726P.
PR	22-JUN-2001; 2001US-300210P.
XX	(COLE-) COLEY PHARM GMBH.
PA	Bauer S, Lipford G, Wagner H;
PI	XX
XX	WPI; 2002-393964/42.
DR	XX
XX	New isolated murine Toll-like receptor (TLR)9, TLR7, TLR8 polypeptides,
PT	useful for identifying species specificity of immunostimulatory nucleic
PT	acid and identifying immunostimulatory nucleic acids -
XX	Disclosure; Page 76; 195pp; English.
PS	XX
XX	The invention relates to isolated murine Toll-like receptors (TLR)9,
CC	TLR7 and TLR8 polypeptides. These polypeptides comprise fully defined,
CC	sequences of 1032, 1050 or 1032 amino acids as given in specification, or
CC	their fragments, where TLR7, TLR7 and TLR8 polypeptides or their
CC	fragments have an amino acid sequence which is identical to human TLR9,
CC	TLR7 or TLR8 polypeptides or their fragment except for at least one amino
CC	acid of a murine TLR polypeptide. The isolated nucleic acids of the
CC	invention are useful for inhibiting TLR9 signalling activity in a cell.
CC	TLR7, TLR8 and TLR9 polypeptides are useful for identifying nucleic acid
CC	molecules which interact with a TLR polypeptide or its fragment. The
CC	TLR7, TLR8 or TLR9 polypeptides are also useful for identifying ISNA. The
CC	TLR7, TLR8 and TLR9 polypeptides are also useful for comparing TLR9
CC	signalling activity of a test compound (that is not a nucleic acid, and
CC	is a polypeptide or a part of a combinatorial library of compounds) with
CC	an ISNA. The TLR7, TLR8 and TLR9 polypeptides are also useful for
CC	identifying species specificity of an ISNA. The isolated nucleic acids of
CC	the invention are useful as probes or primers. This polynucleotide

CC sequence represents DNA relating to the isolated Toll-like receptors of the invention.

XX

SQ Sequence 20 BP; 5 A; 4 C; 7 G; 4 T; 0 other;

Query Match 100.0%; Score 20; DB 24; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.48;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GATTGCGTCAGTCAGAG 20
| | | | | | | | | | | | | | | |
Db 1 GATTGCCTCAGTCAGAG 20

RESULT 5
AAAX76047

ID AAX76047 standard; DNA; 26 BP.

XX

AC AAX76047;

XX

DT 30-JUL-1999 (first entry)

XX

DE CAMP response element oligonucleotide SEQ ID NO:15.

XX

KW CRE; cAMP response element; transcription factor decoy; cis-element;
KW tumour growth inhibitor; palindrome; hairpin; cancer; metabolism;
KW gene transcription regulation; inhibiting proliferation; ds.
XX Synthetic.
OS
XX WO9926634-A1.
PN
XX
PD 03-JUN-1999.
PF 23-NOV-1998; 98WO-US25307.
XX
PR 24-NOV-1997; 97US-0977643.
XX
XX (USSH) US DEPT HEALTH & HUMAN SERVICES.
PA
XX
PI Cho-Chung YS;
XX
XX WPI; 1999-347612/29.
DR
XX
PT Nucleic acids that compete with response elements for transcription factors
PT
XX
XX Example 10; Page 54; 83pp; English.

XX

CC The present invention describes a composition (A) comprising one or more
CC nucleic acids (I) that compete with cAMP (cyclic adenosine monophosphate)
CC response element (CRE) enhancer DNA for binding to transcription factors
CC (TF). (I) are used to regulate gene transcription in cells, in vitro or
CC in vivo, specifically for inhibiting proliferation of cancer cells, but
CC possibly also for regulation of metabolism in hepatitis B and other
CC viruses. HCT-15 human multidrug resistant colon carcinoma cells (2
CC million) were inoculated subcutaneously into the flank of nude mice,
CC then the CRE oligonucleotide 5'-TGAGTTCATGAGTTCATGAGTTCA-3' injected
CC intraperitoneally at doses of 0.1 mg, 5 times per week, once the tumour
CC had reached 30-50 mg. This treatment resulted in over 85% reduction in
CC tumour growth, relative to an untreated control. (I) have high affinity
CC for TF and can inhibit growth of cancer cells without adverse effects on
CC normal cells (contrast use of antisense RNA). The method does not
CC require knowledge of the target gene sequence, only of the response
CC element sequence. The present sequence is used in the exemplification
XX of the present invention.

SQ Sequence 26 BP; 8 A; 4 C; 9 G; 5 T; 0 other;

Query Match 100.0%; Score 20; DB 20; Length 26;
Best Local Similarity 100.0%; Pred. No. 0.49;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCTGACCTCAGAG 20
 Db 4 GATTGCTGACCTCAGAG 23

RESULT 6

AAAT85832
 ID AAT85832 standard; DNA; 27 BP.

XX
 AC AAT85832;

XX 21-NOV-1997 (first entry)

XX CRE oligonucleotide used in gel shift assay.

XX Activating transcription factor 1; ATF1; CREB; recognition sequence;
 KW cyclic AMP responsive element binding protein; inhibition; binding;
 KW proliferation; virus; cancer; HTLV1; leukaemia; antibody; ss.

XX Synthetic.

XX US5641486-A.

XX 24-JUN-1997.

XX 18-MAR-1994; 94US-0210880.

XX 18-MAR-1994; 94US-0210880.

XX (UYNE-) UNIV NEBRASKA.

XX Hinrichs SH, Orten DJ;

XX WPI; 1997-340900/31.

XX Inhibiting replication of cancer cells or viruses - with inhibitor
 PT that binds to peptide sequence of activating transcription factor 1

XX Example 2; Column 6; 17pp; English.

XX This oligonucleotide sequence corresponds to the cyclic AMP binding
 CC element (CRE) to which members of the activating transcription factor 1
 CC (ATF1)-cyclic AMP responsive element binding protein (CREB) family
 CC of protein bind. The sequence was used in a gel shift mobility assay to
 CC identify agents which inhibit the binding of ATF1 to its recognition
 CC sequence. The agents are preferably antibodies, small molecules or
 CC polypeptides, especially the complementarity determining region of
 CC monoclonal antibody Mab4. The agents cause inhibition of transcription
 CC by dissociating ATF1 from its target gene and thus will prevent
 CC proliferation of e.g. a virus or cancer cell, such as HTLV1-mediated
 CC leukaemic cell proliferation.

XX Sequence 27 BP; 8 A; 5 C; 9 G; 5 T; 0 Other;

Query Match 100.0%; Score 20; DB 18; Length 27;
 Best Local Similarity 100.0%; Pred. No. 0.5;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCTGACCTCAGAG 20
 Db 4 GATTGCTGACCTCAGAG 23

RESULT 7

AAV82454
 ID AAV82454 standard; DNA; 27 BP.

XX
 AC AAV82454;

XX 12-APR-1999 (first entry)

XX ATF comp oligonucleotide used in competition analysis.

KW Vascular endothelial growth factor; VEGF; human; hypoxia;
 KW vascular disease; tumour; cancer; angiogenesis; wound healing;
 KW therapy; diagnosis; ds.

XX Synthetic.

OS Homo sapiens.

XX WO9856936-A1.

XX 17-DEC-1998.

XX 10-JUN-1998; 98WO-EP03517.

XX 10-JUN-1997; 97EP-0109418.

XX (PLAC) MAX PLANCK GES FOERDERUNG WISSENSCHAFTEN.

XX Danert A, Plate K, Risau W;

XX WPI; 1999-080911/07.

XX New recombinant DNA - contains sequence that regulates
 PT hypoxia-induced expression, used for, e.g. treatment and diagnosis
 PT of vascular disease

XX Example 6; Page 41; 80pp; English.

XX Oligonucleotides hVEGF, hVEGF 5' DEL, AP1M1 and AP1M2 (see
 CC AAV82449-52), and competitor oligonucleotides AP1 comp, ATF comp
 CC and VL30 (see AAV82453-55) were used in electrophoretic mobility
 CC shift assays to determine which transcription factor(s) bind to
 CC the cis-acting element that is involved in the potentiation of
 CC hypoxia-inducible factor 1 (HIF-1) mediated hypoxic induction
 CC of vascular endothelial growth factor (VEGF) gene regulatory
 CC sequences. Experiments were performed using normoxic or hypoxic
 CC C6 cell nuclear extracts. An AP1 consensus binding site was shown
 CC to compete for DNA-protein complex formation at potentiating
 CC sequences. The invention relates to recombinant DNA molecules
 CC comprising regulatory sequences of the VEGF gene, especially the
 CC 3' untranslated region (see AAV82439) and promoter (see AAV82440),
 CC being capable of modulating hypoxia inducible expression of a
 CC heterologous DNA in vivo. Such recombinant DNA molecules, vectors,
 CC host cells and transgenic animals can be used to identify and
 CC develop compounds and methods for diagnosing, treating, preventing
 CC and/or delaying a vascular or tumour disease.

XX Sequence 27 BP; 8 A; 5 C; 9 G; 5 T; 0 Other;

Query Match 100.0%; Score 20; DB 20; Length 27;
 Best Local Similarity 100.0%; Pred. No. 0.5;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCTGACCTCAGAG 20
 Db 4 GATTGCTGACCTCAGAG 23

RESULT 8

AAV08336

ID AAV08336 standard; DNA; 27 BP.

XX
 AC AAV08336;

XX 04-FEB-1999 (first entry)

XX CRE element coding sequence.

XX ATF1; activating transcription factor 1; inhibitor; gene transcription;
 KW cell proliferation; cancer cell; human; ds.

XX Synthetic.

XX US5844096-A.


```

XX
PD 01-DEC-1998.
XX
PD 20-DEC-1996; 96US-0771411.
XX
PF 20-DEC-1996; 96US-0771411.
XX
PR 18-MAR-1994; 94US-0210880.
XX
PR 20-DEC-1996; 96US-0771411.
XX
XX (UYNE-) UNIV NEBRASKA.
XX
PA Hinrichs SH, Orten DJ;
XX
PI WPI; 1999-044667/04.
XX
DR Inhibitor of activating transcription factor 1 mediated gene
XX PT transcription - useful as anticancer or antiviral agent
XX PT
XX PS Example 2; Column 6; 17pp; English.
XX
PS This sequence represents a CRE element coding sequence. This sequence
CC was used to test the effect of the inhibitory compound of the
CC invention. The inhibitory compound binds to ATrf1 residues 167-181 with
CC sufficient affinity to dissociate ATrf1 from a gene to which it is bound
CC and thereby prevent transcription of the gene. The inhibitory compound
CC and its derivatives are useful for inhibiting the ATrf1-mediated
CC proliferation of cancer cells and viruses, e.g. HTLV I.
XX
SQ Sequence 27 BP; 8 A; 5 C; 9 G; 5 T; 0 other;
Query Match 100.0%; Score 20; DB 20; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.5;.
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCCTGACGTCAGAGAG 20
DB ||||||
4 GATTGCCTGACGTCAGAGAG 23

RESULT 9
AAI70581
ID AAI70581 standard; DNA; 27 BP.
XX
XX AAI70581;
XX
DT 21-JAN-2002 (first entry)
XX
DE Transcription factor CREB consensus oligonucleotide.
XX
KW Transcription factor; CREB; screening; detection; quantification;
KW probe; ds.
XX
OS Synthetic.
XX
PN EP1136567-A1.
XX
PD 26-SEP-2001.
XX
PF 24-MAR-2000; 2000EP-0870057.
XX
PR 24-MAR-2000; 2000EP-0870057.
XX
XX (ADAR-) ADVANCED ARRAY TECHNOLOGIES SA.
XX
PI Remacle J, Renard P, Art M;
XX
XX WPI; 2001-640391/74.
XX
PT Screening, detecting or quantifying transcriptional factors in a
PT biological sample comprises contacting the transcriptional factor with
PT a double-stranded DNA sequence bound to an insoluble solid support
XX
XX Example 4; Page 8; 20pp; English.

```

```
XX SQ Sequence 27 BP; 8 A; 5 C; 9 G; 5 T; 0 other;
Query Match 100.0%; Score 20; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.5;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCGCTGACGTCAGAGAG 20
Db 4 GATTGCGCTGACGTCAGAGAG 23

RESULT 11
AAH77397/c
ID AAH77397 standard; DNA; 27 BP.
XX AC AAH77397;
XX DT 05-NOV-2001 (first entry)
XX DE Cyclic AMP response element CRE consensus oligonucleotide probe #2.
XX KW Cancer; human; dithiocarbamate derivative; anticancer agent; probe; ss.
XX OS Unidentified.
XX PN US2001016600-A1.
XX PD 23-AUG-2001.
XX PF 12-DEC-2000; 2000US-0735205.
XX PR 08-SEP-1998; 98US-0099390.
XX PR 08-SEP-1999; 99US-0392122.
XX PR 05-OCT-2000; 2000US-0679932.
XX PA (KENN/) KENNEDY T P.
XX PI Kennedy TP;
XX PI WPI; 2001-557127/62.
XX DR Treating cancer, asthma and cancer and reducing hypoxic or ischemic
PT damage comprises administering dithiocarbamate thiolate anion or
PT dithiocarbamate thiolate metal complex.
XX PS Disclosure; Page 10; 36pp; English.
XX CC The present invention describes a method of treating cancer, asthma and
CC arthritis and reducing hypoxic or ischaemic damage, involving
CC administering a dithiocarbamate thiolate anion or metal ion complex to
CC the patient. The present sequence is a probe for the cyclic AMP response
CC element CRE, which was described in the exemplification of the invention.
XX SQ Sequence 27 BP; 5 A; 9 C; 5 G; 8 T; 0 other;
Query Match 100.0%; Score 20; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.5;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCGCTGACGTCAGAGAG 20
Db 24 GATTGCGCTGACGTCAGAGAG 5

RESULT 12
AAH87956
ID AAH87956 standard; DNA; 27 BP.
XX AC AAH87956;
XX DT 20-JUL-2001 (first entry)
XX DE Cyclic AMP responsive element CRE consensus oligo for EMSA #2.
XX
```

```
DE XX Cyclic-AMP responsive element CRE consensus oligo for EMSA #1.
XX KW Cyclic-AMP responsive element; electrophoretic mobility shift assay;
KW EMSA; CRE; NF-kappaB; cancer; tetraethyl thiuram disulphide;
KW dithiocarbamate; tumour; metal ion; copper ion; cytokine;
KW ceruloplasmin; anticancer; cytostatic; ss.
XX OS Synthetic.
XX PN WO200117522-A1.
XX PD 15-MAR-2001.
XX PF 15-NOV-1999; 99WO-US27193.
XX PR 08-SEP-1999; 99US-0392122.
XX PA (CHAR-) CHARLOTTE-MECKLENBURG HOSPITAL AUTHORITY.
XX PI Kennedy TP;
XX DR WPI; 2001-281426/29.
XX PT Treating cancer, e.g. melanoma, lung cancer, breast cancer or prostatic
PT carcinoma, comprises administration of a thiuram disulfide optionally
PT in combination with a heavy metal ion, ceruloplasmin or a cytokine e.g.
PT interferon-alpha.
XX PS Disclosure; Page 24; 60pp; English.
XX CC The present invention describes a method for treating established cancer
CC in a mammal. The method comprises administering a thiuram disulfide (I).
CC (I) has anticancer and cytostatic activities. (I) induces apoptosis and
CC G2 cell-cycle arrest, by reducing expression of cyclin A (by preventing
CC binding of transcription factors to DNA regulatory elements involved in
CC control of cyclin A expression). The method can be used to treat cancers,
CC e.g. melanoma, non-small cell lung cancer, small cell lung cancer, renal
CC cancer, colorectal cancer, breast cancer, pancreatic cancer, gastric
CC cancer, bladder cancer, ovarian cancer, uterine cancer, lymphoma and
CC prostate cancer, especially melanoma, lung cancer, breast cancer and
CC prostate carcinoma. The tumour-inhibiting effect of (I) is dependent on
CC heavy metal ions, so administering (I) together with such ions (or with
CC their intracellular carriers, e.g. ceruloplasmin or with serum
CC ceruloplasmin stimulants, e.g. interferon-alpha) will increase the
CC antiproliferative/antineoplastic effect. (I) also potentiates the
CC effect of standard anticancer agents. (I) is already known for treating
CC alcohol abuse (inhibitor of aldehyde dehydrogenase) and is relatively
CC nontoxic and safe. The present sequence represents a cyclic-AMP
CC responsive element CRE consensus oligonucleotide for use in an
CC electrophoretic mobility shift assay (EMSA), which is used in the
CC exemplification of the present invention.
XX SQ Sequence 27 BP; 8 A; 5 C; 9 G; 5 T; 0 other;
Query Match 100.0%; Score 20; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.5;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCGCTGACGTCAGAGAG 20
Db 4 GATTGCGCTGACGTCAGAGAG 23

RESULT 13
AAH87957/c
ID AAH87957 standard; DNA; 27 BP.
XX AC AAH87957;
XX DT 20-JUL-2001 (first entry)
XX DE Cyclic-AMP responsive element CRE consensus oligo for EMSA #2.
XX
```

KW Cyclic-AMP responsive element; electrophoretic mobility shift assay;
 KW EMSA; CRE; NF-kappaB; cancer; tetraethyl thiuram disulphide;
 KW dithiocarbamate; tumour; metal ion; copper ion; cytokine;
 KW ceruloplasmin; anticancer; cytostatic; ss.
 XX Synthetic.
 OS
 XX WO200117522-A1.
 PN
 XX 15-MAR-2001.
 PD
 XX
 PF 15-NOV-1999; 99WO-US27193.
 XX
 PF 08-SEP-1999; 99US-0392122.
 XX
 PR (CHAR-) CHARLOTTE-MECKLENBURG HOSPITAL AUTHORITY.
 PA
 XX Kennedy TP;
 PI
 XX WPI; 2001-281426/29.
 DR
 XX Treating cancer, e.g. melanoma, lung cancer, breast cancer or prostatic
 PT carcinoma, comprises administration of a thiuram disulfide optionally
 PT in combination with a heavy metal ion, ceruloplasmin or a cytokine e.g.
 PT interferon-alpha.
 XX
 PS Disclosure; Page 24; 60pp; English.
 XX
 CC The present invention describes a method for treating established cancer
 CC in a mammal. The method comprises administering a thiuram disulfide (I).
 CC (I) has anticancer and cytostatic activities. (I) induces apoptosis and
 CC G2 cell-cycle arrest, by reducing expression of cyclin A (by preventing
 CC binding of transcription factors to DNA regulatory elements involved in
 CC control of cyclin A expression). The method can be used to treat cancers,
 CC e.g. melanoma, non-small cell lung cancer, small cell lung cancer, renal
 CC cancer, colorectal cancer, breast cancer, pancreatic cancer, gastric
 CC cancer, bladder cancer, ovarian cancer, uterine cancer, lymphoma and
 CC prostate cancer, especially melanoma, lung cancer, breast cancer and
 CC prostate carcinoma. The tumour-inhibiting effect of (I) is dependent on
 CC heavy metal ions, so administering (I) together with such ions (or with
 CC their intracellular carriers, e.g. ceruloplasmin or with serum
 CC ceruloplasmin stimulants, e.g. interferon-alpha) will increase the
 CC antiproliferative/antineoplastic affect. (I) also potentiates the
 CC effect of standard anticancer agents. (I) is already known for treating
 CC alcohol abuse (inhibitor of aldehyde dehydrogenase) and is relatively
 CC nontoxic and safe. The present sequence represents a cyclic-AMP
 CC responsive element CRE consensus oligonucleotide for use in an
 CC electrophoretic mobility shift assay (EMSA), which is used in the
 CC exemplification of the present invention.
 XX
 SQ Sequence 27 BP; 5 A; 9 C; 5 G; 8 T; 0 other;
 Query Match 100.0%; Score 20; DB 22; Length 27;
 Best Local Similarity 100.0%; Pred. No. 0.5;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GATTGCTGACGTCAGAG 20
 ID AAF76267
 XX ||||||||||||||||
 DB 24 GATTGCTGACGTCAGAG 5
 XX
 RESULT 14
 ID AAF76267
 XX AAF76267 standard; DNA; 27 BP.
 AC AAF76267;
 XX
 DT 05-JUN-2001 (first entry)
 XX
 DE CAMP response element (CRE) competitor EMSA probe.
 XX
 KW NF-kappa-B; nuclear factor-kappa-B; CAMP response element; CRE;
 KW nuclear translocation inhibition; heparin; internalisation;

KW NF-kappa-B dependent gene expression inhibition; cytokine;
 KW tumour necrosis factor; TNF; interleukin; IL-1; IL-2; IL-6; IL-8;
 KW interferon-beta; interferon-gamma; tissue factor-1; complement;
 KW inducible nitric oxide synthase; diabetic vascular disease;
 KW heart failure; asthma; sepsis; ischaemic-reperfusion injury;
 KW electrophoretic mobility shift assay; competitor EMSA probe; ds.
 XX
 OS Unidentified.
 XX
 PN WO200119376-A2.
 PD
 XX 22-MAR-2001.
 PF
 XX 12-SEP-2000; 2000WO-US24910.
 XX
 PR 13-SEP-1999; 99US-0395081.
 XX
 PA (CHAR-) CHARLOTTE-MECKLENBURG HOSPITAL AUTHORITY.
 PI
 XX Kennedy TP;
 DR
 XX WPI; 2001-244698/25.
 XX
 CC Inhibiting NF-kappa-B activity, useful for treating e.g. diabetic
 PT vascular disease, heart failure, asthma and sepsis, comprises
 PT administering heparin to cells in patient to inhibit translocation of
 PT NF-kappa-B from cytoplasm to nucleus.
 XX
 PS Examples; Page 22; 58pp; English.
 XX
 CC The invention relates to a method of inhibiting nuclear factor-kappa-B
 CC (NF-kappa-B) activity in a patient, comprising the administration of
 CC heparin to the cells in the patient, such that the heparin is
 CC internalised into the cytoplasm of cells in the patient. The invention
 CC is based on the discovery that heparin is able to block the
 CC translocation of NF-kappa-B from the cytoplasm to the nucleus. This in
 CC turn inhibits NF-kappa-B dependent gene expression. Such NF-kappa-B
 CC dependent genes include genes encoding cytokines such as tumour necrosis
 CC factor (TNF), IL-1 (interleukin-1), IL-2, IL-6, IL-8, interferon-beta,
 CC interferon-gamma, tissue factor-1, complement and inducible nitric
 CC oxide synthase. The method of the invention is used for treating or
 CC preventing diabetic vascular disease, heart failure, asthma, sepsis and
 CC ischaemic-reperfusion injury. Heparin may be administered in combination
 CC with other active agents that treat or prevent another disease or
 CC symptom in the patient, e.g., antiviral agents, antibiotics, antifungal
 CC agents and antiinflammatory agents. The method of the invention offers
 CC significant advantages over prior art treatments for the above
 CC conditions. Heparin is relatively non-toxic and safe, and should not
 CC produce the side effects such as hypertension, glucose intolerance
 CC and bone demineralisation that are encountered with the use of
 CC glucocorticoids for blocking the NF-kappa-B nuclear translocation.
 CC Additionally, heparin is readily available and easily used. Sequences
 CC AAF76266-AAF76267 represents EMSA (electrophoretic mobility shift assay)
 CC probes used to measure the effect of heparin on NF-kappa-B nuclear
 CC translocation. EMSA probe AAF76266 comprises a consensus NF-kappa-B
 CC response element, and EMSA competitor probe AAF76267 comprises a
 CC CAMP response element (CRE).
 XX
 SQ Sequence 27 BP; 8 A; 5 C; 9 G; 5 T; 0 other;

Query Match 100.0%; Score 20; DB 22; Length 27;
 Best Local Similarity 100.0%; Pred. No. 0.5;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GATTGCTGACGTCAGAG 20
 ID ABA92274
 XX ||||||||||||||||
 DB 4 GATTGCTGACGTCAGAG 23
 XX
 RESULT 15
 ID ABA92274
 XX ABA92274 standard; DNA; 27 BP.
 XX

us-09-355-254f-17.rni

Thu Dec 12 07:53:51 2002

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OM nucleic - nucleic search, using sw model
Run on: December 12, 2002, 01:05:40 : Search time 22.2464 Seconds
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Title: US-09-355-254F-17
Perfect score: 20
Sequence: 1 tatgcataattcctgaagt 20

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 153338381 residues 687286
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Listing first 45 summaries

Database : Issued Patents, NA: *
1: /cgn2_6/ptodata/1/ina/5A_COMB.seq: *
2: /cgn2_6/ptodata/1/ina/5B_COMB.seq: *
3: /cgn2_6/ptodata/1/ina/6A_COMB.seq: *
4: /cgn2_6/ptodata/1/ina/6B_COMB.seq: *
5: /cgn2_6/ptodata/1/ina/PTUS_COMB.seq: *
6: /cgn2_6/ptodata/1/ina/backfiles1.seq: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	16	80.0	19	1	US-08-410-780A-75
2	16	80.0	19	1	US-08-411-020-40
3	16	80.0	19	5	PCT-US95-04511-75
4	15	75.0	17	1	US-08-369-796-21
5	15	75.0	17	2	US-08-852-091-21
6	15	75.0	17	5	PCT-US95-17025-21
7	15	75.0	19	1	US-08-410-780A-76
8	15	75.0	19	1	US-08-411-020-41
9	15	75.0	19	5	PCT-US95-04511-76
10	14.8	74.0	47	1	US-08-171-389-37
11	14.8	74.0	47	2	US-08-123-936-37
12	14.8	74.0	47	3	US-08-475-228A-37
13	14.8	74.0	47	3	US-08-482-080A-37
14	14.8	74.0	47	4	US-09-354-947-37
15	14.8	74.0	47	5	PCT-US93-12388-37
16	14.8	74.0	50	1	US-08-171-389-437
17	14.8	74.0	50	1	US-08-123-936-437
18	14.8	74.0	50	1	US-08-171-389-437
19	14.8	74.0	50	1	US-08-123-936-437
20	14.8	74.0	50	2	US-08-475-228A-437
21	14.8	74.0	50	2	US-08-475-228A-437
22	14.8	74.0	50	3	US-08-482-080A-437
23	14.8	74.0	50	3	US-08-482-080A-437
24	14.8	74.0	50	4	US-09-354-947-437
25	14.8	74.0	50	4	US-09-354-947-437
26	14.8	74.0	50	5	PCT-US93-12388-437
27	14.8	74.0	50	5	PCT-US93-12388-437

28	14.4	72.0	19	1	US-08-410-780A-23	Sequence 23, Appl
29	14.4	72.0	19	1	US-08-410-780A-25	Sequence 25, Appl
30	14.4	72.0	19	1	US-08-410-780A-33	Sequence 33, Appl
31	14.4	72.0	19	1	US-08-410-780A-49	Sequence 49, Appl
32	14.4	72.0	19	1	US-08-410-780A-51	Sequence 51, Appl
33	14.4	72.0	19	1	US-08-410-780A-55	Sequence 55, Appl
34	14.4	72.0	19	1	US-08-411-020-42	Sequence 42, Appl
35	14.4	72.0	19	5	PCT-US95-04511-23	Sequence 23, Appl
36	14.4	72.0	19	5	PCT-US95-04511-25	Sequence 25, Appl
37	14.4	72.0	19	5	PCT-US95-04511-33	Sequence 33, Appl
38	14.4	72.0	19	5	PCT-US95-04511-49	Sequence 49, Appl
39	14.4	72.0	19	5	PCT-US95-04511-51	Sequence 51, Appl
40	14.4	72.0	19	5	PCT-US95-04511-55	Sequence 55, Appl
41	14	70.0	20	4	US-09-489-868A-31	Sequence 31, Appl
42	13.4	67.0	19	1	US-08-410-780A-24	Sequence 24, Appl
43	13.4	67.0	19	1	US-08-410-780A-26	Sequence 26, Appl
44	13.4	67.0	19	1	US-08-410-780A-34	Sequence 34, Appl
45	13.4	67.0	19	1	US-08-410-780A-50	Sequence 50, Appl

ALIGNMENTS

RESULT 1
US-08-410-780A-75
; Sequence 75, Application US/08410780A
; Patent No. 5707803
; GENERAL INFORMATION:
; APPLICANT: LAMB, I. PETER
; APPLICANT: SEIDEL, H. MARTI
; TITLE OF INVENTION: DNA REGULATORY ELEMENTS RESPONSIVE TO
; TITLE OF INVENTION: CYTOKINES AND METHODS FOR THEIR USE
; NUMBER OF SEQUENCES: 77
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: LIGAND PHARMACEUTICALS INCORPORATED
; STREET: 9393 TOWNE CENTRE DRIVE
; CITY: SAN DIEGO
; STATE: CALIFORNIA
; COUNTRY: US
; ZIP: 92121

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/410,780A
FILING DATE: 27-MAR-1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/228,934
FILING DATE: 14-APR-1994
ATTORNEY/AGENT INFORMATION:
NAME: JURGENSEN, THOMAS E
REGISTRATION NUMBER: 34,195
REFERENCE/DOCKET NUMBER: 016-0017A.US
TELEPHONE: (619) 550-7675
TELEFAX: (619) 535-3906
INFORMATION FOR SEQ ID NO: 75:
SEQUENCE CHARACTERISTICS:
LENGTH: 19 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "OTHER NUCLEIC ACID, SYNTHETIC DNA"

Query Match 80.0%; Score 16; DB 1; Length 19;
Best Local Similarity 100.0%; Pred No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 CATATTCCTGTAAGTG 20
DB 4 CATATTCCTGTAAGTG 19

RESULT 2

US-08-411-020-40
; Sequence 40, Application US/08411020
; Patent No. 5712094
; GENERAL INFORMATION:
; APPLICANT: SEIDEL, H. MARTI
; APPLICANT: LAMB, I. PETER
; APPLICANT: CHAN, SHIN-SHAY TIAN
; TITLE OF INVENTION: METHODS AND ASSOCIATED REAGENTS FOR
; TITLE OF INVENTION: DETECTING MODULATORS OF CYTOKINE ACTION
; NUMBER OF SEQUENCES: 59
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Ligand Pharmaceuticals Incorporated
; STREET: 9393 Towne Centre Drive
; CITY: San Diego
; STATE: California
; COUNTRY: US
; ZIP: 92121

COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/411,020
; FILING DATE: 27-MAR-1995
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Jurgensen, Thomas E.
; REGISTRATION NUMBER: 34,195
; REFERENCE/DOCKET NUMBER: 016-0030.US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 550-7675
; TELEFAX: (619) 535-3906
; INFORMATION FOR SEQ ID NO: 40:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 19 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: /desc = "OTHER NUCLEIC ACID,
; DESCRIPTION: SYNTHETIC DNA"
US-08-411-020-40

Query Match 80.0%; Score 16; DB 1; Length 19;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 CATATTCCTGTAAGTG 20
DB 4 CATATTCCTGTAAGTG 19

RESULT 3

PCT-US95-04511-75
; Sequence 75, Application PC/TUS9504511
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: DNA REGULATORY ELEMENTS RESPONSIVE TO
; TITLE OF INVENTION: CYTOKINES AND METHODS FOR THEIR USE
; NUMBER OF SEQUENCES: 76
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)

; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/04511
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/228,934
; FILING DATE: 14-APR-1994
; INFORMATION FOR SEQ ID NO: 75:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 19 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: /desc = "OTHER NUCLEIC ACID,
; DESCRIPTION: SYNTHETIC DNA"
PCT-US95-04511-75

Query Match 80.0%; Score 16; DB 5; Length 19;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 CATATTCCTGTAAGTG 20
DB 4 CATATTCCTGTAAGTG 19

RESULT 4

US-08-369-796-21
; Sequence 21, Application US/08369796
; Patent No. 5716622
; GENERAL INFORMATION:
; APPLICANT: James E. Darnell, Jr.
; APPLICANT: Zilong Wen
; APPLICANT: Curt M. Horvath
; APPLICANT: Zhong Zhong
; TITLE OF INVENTION: FUNCTIONALLY ACTIVE REGIONS OF SIGNAL
; TITLE OF INVENTION: TRANSDUCER AND ACTIVATOR OF TRANSCRIPTION (STAT) PROTEINS
; NUMBER OF SEQUENCES: 39
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Klauber & Jackson
; STREET: 411 Hackensack Avenue
; CITY: Hackensack
; STATE: New Jersey
; COUNTRY: USA
; ZIP: 07601

COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/369,796
; FILING DATE: 06-JAN-1995
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Jackson Esq., David A.
; REGISTRATION NUMBER: 26,742
; REFERENCE/DOCKET NUMBER: 600-1-116
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 201 487-5800
; TELEFAX: 201 343-1684
; TELEX: 133521
; INFORMATION FOR SEQ ID NO: 21:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 17 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA synthetic probe
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
US-08-369-796-21

```
Query Match 75.0%; Score 15; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 47;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATTCCTGTAAGTG 20
Db 1 ATATTCCTGTAAGTG 15

RESULT 5
US-08-852-091-21
; Sequence 21, Application US/08852091
; Patent No. 5863228
; GENERAL INFORMATION:
; APPLICANT: James E. Darnell, Jr.
; APPLICANT: Zilong Wen
; APPLICANT: Curt M. Horvath
; APPLICANT: Zhong Zhong
; TITLE OF INVENTION: FUNCTIONALLY ACTIVE REGIONS OF SIGNAL
; TITLE OF INVENTION: TRANSDUCER AND ACTIVATOR OF TRANSCRIPTION (STAT) PROTEINS
; NUMBER OF SEQUENCES: 39
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Klauber & Jackson
; STREET: 411 Hackensack Avenue
; CITY: Hackensack
; STATE: New Jersey
; COUNTRY: USA
; ZIP: 07601
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION NUMBER:
; FILING DATE: 06-MAY-1997
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/852,091
; FILING DATE: 06-MAY-1997
; CLASSIFICATION: 424
; APPLICATION DATA:
; FILING DATE: 06-JAN-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Jackson Esq., David A.
; REGISTRATION NUMBER: 26,742
; REFERENCE/DOCKET NUMBER: 600-1-116
; TELEPHONE: 201 487-5800
; TELEFAX: 201 343-1684
; TELEX: 133521
; INFORMATION FOR SEQ ID NO: 21:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 17 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA synthetic probe
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; US-08-852-091-21

Query Match 75.0%; Score 15; DB 2; Length 17;
Best Local Similarity 100.0%; Pred. No. 47;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATTCCTGTAAGTG 20
Db 1 ATATTCCTGTAAGTG 15

RESULT 6
PCT-US95-17025-21
; Sequence 21, Application PC/TUS9517025
; GENERAL INFORMATION:
```

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; APPLICANT: James E. Darnell, Jr.
; APPLICANT: Zilong Wen
; APPLICANT: Curt M. Horvath
; APPLICANT: Zhong Zhong
; TITLE OF INVENTION: FUNCTIONALLY ACTIVE REGIONS OF SIGNAL
; TITLE OF INVENTION: TRANSDUCER AND ACTIVATOR OF TRANSCRIPTION (STAT) PROTEINS
; NUMBER OF SEQUENCES: 39
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Klauber & Jackson
; STREET: 411 Hackensack Avenue
; CITY: Hackensack
; STATE: New Jersey
; COUNTRY: USA
; ZIP: 07601
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION NUMBER: PCT/US95/17025
; FILING DATE: 28-DEC-1995
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/369,796
; FILING DATE: 06-JAN-1995
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Jackson Esq., David A.
; REGISTRATION NUMBER: 26,742
; REFERENCE/DOCKET NUMBER: 600-1-116
; TELEPHONE: 201 487-5800
; TELEFAX: 201 343-1684
; TELEX: 133521
; INFORMATION FOR SEQ ID NO: 21:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 17 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA synthetic probe
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; PCT-US95-17025-21

Query Match 75.0%; Score 15; DB 5; Length 17;
Best Local Similarity 100.0%; Pred. No. 47;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATTCCTGTAAGTG 20
Db 1 ATATTCCTGTAAGTG 15

RESULT 7
US-08-410-780A-76/c
; Sequence 76, Application US/08410780A
; Patent No. 5707803
; GENERAL INFORMATION:
; APPLICANT: LAMB, I. PETER
; APPLICANT: SEIDEL, H. MARTI
; TITLE OF INVENTION: DNA REGULATORY ELEMENTS RESPONSIVE TO
; TITLE OF INVENTION: CYTOKINES AND METHODS FOR THEIR USE
; NUMBER OF SEQUENCES: 77
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: LIGAND PHARMACEUTICALS INCORPORATED
; STREET: 9393 TOWNE CENTRE DRIVE
; CITY: SAN DIEGO
; STATE: CALIFORNIA
; COUNTRY: US
; ZIP: 92121
; COMPUTER READABLE FORM:
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;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: PatentIn Release #1.0, Version #1.30
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/410,780A
;; FILING DATE: 27-MAR-1995
;; CLASSIFICATION: 435
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 08/228,934
;; FILING DATE: 14-APR-1994
;; ATTORNEY/AGENT INFORMATION:
;; NAME: JURGENSEN, THOMAS E
;; REGISTRATION NUMBER: 34,195
;; REFERENCE/DOCKET NUMBER: 016-0017A.US
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: (619) 550-7675
;; TELEFAX: (619) 535-3906
;; INFORMATION FOR SEQ ID NO: 76:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 19 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: other nucleic acid
;; DESCRIPTION: /desc = "OTHER NUCLEIC ACID,
;; SYNTHETIC DNA"
US-08-410-780A-76

Query Match 75.0%; Score 15; DB 1; Length 19;
Best Local Similarity 100.0%; Pred. No. 48;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATTCCTGTAAGTG 20
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Db 19 ATATTCCTGTAAGTG 5

RESULT 8
US-08-411-020-41/c
;; Sequence 41, Application US/08411020
;; Patent No. 5712094
;; GENERAL INFORMATION:
;; APPLICANT: SEIDEL, H. MARTI
;; APPLICANT: LAMB, I. PETER
;; APPLICANT: CHAN, SHIN-SHAY TIAN
;; TITLE OF INVENTION: METHODS AND ASSOCIATED REAGENTS FOR
;; DETECTING MODULATORS OF CYTOKINE ACTION
;; NUMBER OF SEQUENCES: 59
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: Ligand Pharmaceuticals Incorporated
;; STREET: 9393 Towne Centre Drive
;; CITY: San Diego
;; STATE: California
;; COUNTRY: US
;; ZIP: 92121
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: PatentIn Release #1.0, Version #1.30
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/411,020
;; FILING DATE: 27-MAR-1995
;; CLASSIFICATION: 435
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Jurgensen, Thomas E.
;; REGISTRATION NUMBER: 34,195
;; REFERENCE/DOCKET NUMBER: 016-0030.US
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: (619) 550-7675
;; TELEFAX: (619) 535-3906
;; INFORMATION FOR SEQ ID NO: 41:

;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 19 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: other nucleic acid
;; DESCRIPTION: /desc = "OTHER NUCLEIC ACID,
;; SYNTHETIC DNA"
US-08-411-020-41

Query Match 75.0%; Score 15; DB 1; Length 19;
Best Local Similarity 100.0%; Pred. No. 48;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATTCCTGTAAGTG 20
|||||
Db 19 ATATTCCTGTAAGTG 5

RESULT 9
PCT-US95-04511-76/c
;; Sequence 76, Application PC/TUS9504511
;; GENERAL INFORMATION:
;; APPLICANT:
;; TITLE OF INVENTION: DNA REGULATORY ELEMENTS RESPONSIVE TO
;; CYTOKINES AND METHODS FOR THEIR USE
;; NUMBER OF SEQUENCES: 76
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: PCT/US95/04511
;; FILING DATE:
;; CLASSIFICATION:
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 08/228,934
;; FILING DATE: 14-APR-1994
;; INFORMATION FOR SEQ ID NO: 76:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 19 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: other nucleic acid
;; DESCRIPTION: /desc = "OTHER NUCLEIC ACID,
;; SYNTHETIC DNA"
PCT-US95-04511-76

Query Match 75.0%; Score 15; DB 5; Length 19;
Best Local Similarity 100.0%; Pred. No. 48;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATTCCTGTAAGTG 20
|||||
Db 19 ATATTCCTGTAAGTG 5

RESULT 10
US-08-171-389-37/c
;; Sequence 37, Application US/08171389
;; Patent No. 5578444
;; GENERAL INFORMATION:
;; APPLICANT: Edwards, Cynthia A.
;; APPLICANT: Cantor, Charles R.
;; APPLICANT: Andrews, Beth M.
;; APPLICANT: Turin, Lisa M.
;; APPLICANT: Fry, Kirk E.
;; TITLE OF INVENTION: Sequence-Directed DNA Binding
;; MOLECULES, COMPOSITIONS AND METHODS
;; NUMBER OF SEQUENCES: 641
;; CORRESPONDENCE ADDRESS:

ADDRESSEE: Genelabs Technologies, Inc.
STREET: 505 Penobscot Drive
CITY: Redwood City
STATE: CA
COUNTRY: USA
ZIP: 94063
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/171,389
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/123,936
FILING DATE: 17-SEP-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/996,783
FILING DATE: 23-DEC-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/723,618
FILING DATE: 27-JUN-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/081,070
FILING DATE: 22-JUN-1993
ATTORNEY/AGENT INFORMATION:
NAME: Fabian, Gary R.
REGISTRATION NUMBER: 33,875
REFERENCE/DOCKET NUMBER: 4600-0175/G19P3
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 324-0880
TELEFAX: (415) 324-0960
INFORMATION FOR SEQ ID NO: 37:
SEQUENCE CHARACTERISTICS:
LENGTH: 47 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ORIGINAL SOURCE:
INDIVIDUAL ISOLATE: Human alpha-amylase gene
US-08-171-389-37

Query Match 74.0%; Score 14.8; DB 1; Length 47;
Best Local Similarity 88.9%; Pred. No. 67;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18
||| |||||||||
Db 28 TATTATATTCCTGTAAG 11

RESULT 11
US-08-123-936-37/c
Sequence 37, Application US/08123936
Patent No. 5726014
GENERAL INFORMATION:
APPLICANT: Edwards, Cynthia A.
APPLICANT: Cantor, Charles R.
APPLICANT: Andrews, Beth M.
APPLICANT: Turin, Lisa M.
TITLE OF INVENTION: Screening Assay for the Detection of
NUMBER OF SEQUENCES: 640
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genelabs Technologies, Inc.
STREET: 505 Penobscot Drive
CITY: Redwood City
STATE: CA
COUNTRY: USA

ZIP: 94063
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/123,936
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/996,783
FILING DATE: 23-DEC-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/723,618
FILING DATE: 27-JUN-1991
ATTORNEY/AGENT INFORMATION:
NAME: Fabian, Gary R.
REGISTRATION NUMBER: 33,875
REFERENCE/DOCKET NUMBER: 4600-0075.32/G19P2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 324-0880
TELEFAX: (415) 324-0960
INFORMATION FOR SEQ ID NO: 37:
SEQUENCE CHARACTERISTICS:
LENGTH: 47 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ORIGINAL SOURCE:
INDIVIDUAL ISOLATE: Human alpha-amylase gene
US-08-123-936-37

Query Match 74.0%; Score 14.8; DB 1; Length 47;
Best Local Similarity 88.9%; Pred. No. 67;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18
||| |||||||||
Db 28 TATTATATTCCTGTAAG 11

RESULT 12
US-08-475-228A-37/c
Sequence 37, Application US/08475228A
Patent No. 5869241
GENERAL INFORMATION:
APPLICANT: Edwards, Cynthia A.
APPLICANT: Cantor, Charles R.
APPLICANT: Andrews, Beth M.
APPLICANT: Turin, Lisa M.
APPLICANT: Fry, Kirk E.
TITLE OF INVENTION: Sequence-Directed DNA Binding
NUMBER OF SEQUENCES: 664
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genelabs Technologies, Inc.
STREET: 505 Penobscot Drive
CITY: Redwood City
STATE: CA
COUNTRY: USA
ZIP: 94063
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/475,228A
FILING DATE: 06-JUN-1995
PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/123,936
FILING DATE: 17-SEP-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/996,783
FILING DATE: 23-DEC-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/723,618
FILING DATE: 27-JUN-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/081,070
FILING DATE: 22-JUN-1993
ATTORNEY/AGENT INFORMATION:
NAME: Stratford, Carol A.
REGISTRATION NUMBER: 34,444
REFERENCE/DOCKET NUMBER: 4600-0175.21/G19P3D2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 324-0880
TELEFAX: (415) 324-0960
INFORMATION FOR SEQ ID NO: 37:
SEQUENCE CHARACTERISTICS:
LENGTH: 47 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ORIGINAL SOURCE:
INDIVIDUAL ISOLATE: Human alpha-amylase gene
US-08-475-228A-37

Query Match 74.0%; Score 14.8; DB 2; Length 47;
Best Local Similarity 88.9%; Pred. No. 67;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18
||| |||||
Db 28 TATTATATTCCTGTAAG 11

RESULT 13
US-08-482-080A-37/C
Sequence 37, Application US/08482080A
Patent No. 6010849
GENERAL INFORMATION:
APPLICANT: Edwards, Cynthia A.
APPLICANT: Cantor, Charles R.
APPLICANT: Andrews, Beth M.
APPLICANT: Turin, Lisa M.
APPLICANT: Fry, Kirk E.
TITLE OF INVENTION: Sequence-Directed DNA Binding
Molecules, Compositions and Methods
NUMBER OF SEQUENCES: 664
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genelabs Technologies, Inc.
STREET: 505 Penobscot Drive
CITY: Redwood City
STATE: CA
COUNTRY: USA
ZIP: 94063
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/482,080A
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/171,389
FILING DATE: 20-DEC-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/123,936
FILING DATE: 17-SEP-1993

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/996,783
FILING DATE: 23-DEC-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/723,618
FILING DATE: 27-JUN-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/081,070
FILING DATE: 22-JUN-1993
ATTORNEY/AGENT INFORMATION:
NAME: Brady, John F.
REGISTRATION NUMBER: 39,118
REFERENCE/DOCKET NUMBER: 4600-0175.20/G19P3D1
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 324-0880
TELEFAX: (650) 324-0960
INFORMATION FOR SEQ ID NO: 37:
SEQUENCE CHARACTERISTICS:
LENGTH: 47 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ORIGINAL SOURCE:
INDIVIDUAL ISOLATE: Human alpha-amylase gene
US-08-482-080A-37

Query Match 74.0%; Score 14.8; DB 3; Length 47;
Best Local Similarity 88.9%; Pred. No. 67;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18
||| |||||
Db 28 TATTATATTCCTGTAAG 11

RESULT 14
US-09-354-947-37/C
Sequence 37, Application US/09354947
Patent No. 6384208
GENERAL INFORMATION:
APPLICANT: Edwards, Cynthia A.
APPLICANT: Cantor, Charles R.
APPLICANT: Andrews, Beth M.
APPLICANT: Turin, Lisa M.
APPLICANT: Fry, Kirk E.
TITLE OF INVENTION: Sequence-Directed DNA Binding
Molecules, Compositions and Methods
NUMBER OF SEQUENCES: 664
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genelabs Technologies, Inc.
STREET: 505 Penobscot Drive
CITY: Redwood City
STATE: CA
COUNTRY: USA
ZIP: 94063
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/354,947
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/482,080
FILING DATE: 07-JUN-1995
APPLICATION NUMBER: US 08/171,389
FILING DATE: 20-DEC-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/123,936
FILING DATE: 17-SEP-1993

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/996,783
FILING DATE: 23-DEC-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/723,618
FILING DATE: 27-JUN-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/081,070
FILING DATE: 22-JUN-1993
ATTORNEY/AGENT INFORMATION:
NAME: Brady, John F.
REGISTRATION NUMBER: 39,118
REFERENCE/DOCKET NUMBER: 4600-0175.20/G19P3D1
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 324-0880
TELEFAX: (650) 324-0960
INFORMATION FOR SEQ ID NO: 37:
SEQUENCE CHARACTERISTICS:
LENGTH: 47 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ORIGINAL SOURCE:
INDIVIDUAL ISOLATE: Human alpha-amylase gene
US-09-354-947-37

Query Match 74.0%; Score 14.8; DB 4; Length 47;
Best Local Similarity 88.9%; Pred. No. 67;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18
||| |||||
Db 28 TATTATATTCCTGTAAG 11

RESULT 15
PCT-US93-12388-37/c
Sequence 37, Application PC/TUS9312388
GENERAL INFORMATION:
APPLICANT:
TITLE OF INVENTION: Sequence-Directed DNA Binding
TITLE OF INVENTION: Molecules, Compositions and Methods
NUMBER OF SEQUENCES: 641
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genelabs Technologies, Inc.
STREET: 505 Penobscot Drive
CITY: Redwood City
STATE: CA
COUNTRY: USA
ZIP: 94063
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US93/12388
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/123,936
FILING DATE: 17-SEP-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/996,783
FILING DATE: 23-DEC-1992
ATTORNEY/AGENT INFORMATION:
NAME: Fabian, Gary R.
REGISTRATION NUMBER: 33,875
REFERENCE/DOCKET NUMBER: 4600-0175.41/G19PCT2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 324-0880

TELEFAX: (415) 324-0960
INFORMATION FOR SEQ ID NO: 37:
SEQUENCE CHARACTERISTICS:
LENGTH: 47 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ORIGINAL SOURCE:
INDIVIDUAL ISOLATE: Human alpha-amylase gene
PCT-US93-12388-37
Query Match 74.0%; Score 14.8; DB 5; Length 47;
Best Local Similarity 88.9%; Pred. No. 67;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 TATGCATATTCCTGTAAG 18
||| |||||
Db 28 TATTATATTCCTGTAAG 11

Search completed: December 12, 2002, 01:41:54
Job time : 22.2464 secs

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OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:04:50 ; Search time 102.319 Seconds
(without alignments)
440.192 Million cell updates/sec

Title: US-09-355-254F-17

Perfect score: 20

Sequence: 1 tatgatattcttgaagtg 20

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 112599159 residues

Total number of hits satisfying chosen parameters: 2390332

Minimum DB seq length: 0

Maximum DB seq length: 100

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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21: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2000.DAT.*
22: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
23: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
24: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20	100.0	20	19	AAV46009
2	20	100.0	20	24	AA139193
3	19	95.0	24	24	ABN80818
4	16	80.0	19	17	AA141604
5	15	75.0	17	17	AA131283
6	14.8	74.0	47	15	AAQ69287
7	14.8	74.0	47	18	AA163749
8	14.8	74.0	47	20	AA17037
9	14.8	74.0	47	24	ABR82528

C 10	14.8	74.0	50	15	AAQ69686	Human pancreatic a
C 11	14.8	74.0	50	15	AAQ69687	Human pancreatic a
C 12	14.8	74.0	50	18	AA164148	Human pancreatic a
C 13	14.8	74.0	50	18	AA164149	Human pancreatic a
C 14	14.8	74.0	50	20	AA117436	Test sequence from
C 15	14.8	74.0	50	20	AA117437	DNA binding molecu
C 16	14.8	74.0	50	24	ABK82927	DNA binding molecu
C 17	14.8	74.0	50	24	ABK82928	Oligonucleotide co
C 18	14.4	72.0	19	17	AA141605	Human cot oncogene
C 19	14	70.0	20	22	AA111323	Primer 2 for pMAL-
C 20	14	70.0	27	15	AA114907	C2P-3 Gene primer.
C 21	14	70.0	27	15	AAQ55426	Single nucleotide
C 22	13.8	69.0	22	21	AA171335	Humanised anti-Fas
C 23	13.6	68.0	50	19	AAV66651	Humanised anti-Fas
C 24	13.6	68.0	50	21	AA178342	PEBP2 alpha A gene
C 25	13.4	67.0	21	20	AA133318	Oligonucleotide ad
C 26	13.4	67.0	24	24	ABQ01109	Oligonucleotide ad
C 27	13.4	67.0	24	24	ABQ06151	Oligonucleotide ad
C 28	13.4	67.0	24	24	ABQ06192	Oligonucleotide ad
C 29	13.4	67.0	25	22	AA139126	SNP specific lower
C 30	13.4	67.0	51	22	AA179532	Human silent SNP c
C 31	13.4	67.0	60	14	AAQ51030	5' fragment of the
C 32	13.4	67.0	60	24	ABN39544	Human spliced tran
C 33	13.2	66.0	21	22	AA187028	Sequencing primer
C 34	13.2	66.0	31	20	AA138743	Human genomic DNA
C 35	13.2	66.0	60	24	ABN36070	Human spliced tran
C 36	13.2	66.0	65	24	ABN29969	Rat spliced trans
C 37	13.2	66.0	99	21	AA242637	Human 5' EST isola
C 38	13	65.0	13	16	AA102747	Ly6 GAS regulatory
C 39	13	65.0	47	21	AA266670	Human map-related
C 40	13	65.0	65	24	ABN52711	Mouse spliced tran
C 41	12.8	64.0	23	20	AA152734	Human genome biall
C 42	12.8	64.0	36	13	AAQ31180	PCR primer #2 for
C 43	12.8	64.0	41	24	ABK48179	Citrate synthase 1
C 44	12.8	64.0	47	20	AA152584	Human genome biall
C 45	12.8	64.0	47	20	AA152534	Human genome biall

ALIGNMENTS

RESULT 1
AAV46009
ID AAV46009 standard; DNA; 20 BP.

XX AAV46009;

XX 16-OCT-1998 (first entry)

DE Immune adjuvant STAT1.

XX Immune system; adjuvant; vaccine; cancer; prophylactic; pathogenicity;
modulator; tolerance; regulator; helper cell; antigen; immunoglobulin;
XX Ig class; autoimmune response; T-cell; B-cell; tumour; ss.

OS Class Bacteria.

XX EP855184-A1.

XX 29-JUL-1998.

XX 23-JAN-1997; 97EP-0101019.

XX 23-JAN-1997; 97EP-0101019.

XX (HEG/) HEG K.

XX (LIPF/) LIPFORD G B.

XX (WAGN/) WAGNER H.

XX Heeg K, Lipford GB, Wagner H;

XX WPI; 1998-389630/34.

XX

PT Antigenic composition comprises polynucleotide fragment and antigen
PT used as vaccine to treat or prevent e.g. cancer or pathogen
PT infections and to modulate immune response e.g. tolerance break and
PT regulation of TH1/TH2 cells
XX
XX Example 5; Page 9; 28pp; English.
PS
XX
CC AAV5993-V46019 are fragments of bacterial polynucleotides which are
CC used as immune adjuvants for inclusion into vaccines to treat cancer and
CC for prophylaxis and/or treatment of conditions caused by pathogenic
CC micro-organisms. The polynucleotide is used for modulation of an immune
CC response and the modulation is selected from the group break of
CC tolerance, regulation of TH1/TH2 helper cell responses, switch of Ig
CC classes, treatment of autoimmune responses and induction of tolerances.
CC DNA oligomers are used to enhance the reactivity of immune cells to
CC viral, bacterial and parasitic antigens, to break tolerance in anergic T
CC and B cells e.g. against tumour antigens, as adjuvants in vaccination
CC against tumour-defined antigens and immunostimulatory substances in an
CC immune response against tumours and to suppress immune reactions of the
CC innate and acquired immune system. The composition is inexpensive and
CC stable and does not cause lethal shock, which happens with prior art
CC bacterial sequences.
XX
SQ Sequence 20 BP; 5 A; 3 C; 4 G; 8 T; 0 other;
XX
XX Query Match 100.0%; Score 20; DB 19; Length 20;
XX Best Local Similarity 100.0%; Pred. No. 1.3;
XX Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY 1 TATGCATATTCCTGTAAGTG 20
Db | | | | | | | | | | | | | | | | | | | |
1 TATGCATATTCCTGTAAGTG 20
XX
XX
XX RESULT 2
XX AAL39193
XX ID AAL39193 standard; DNA; 20 BP.
XX AC AAL39193;
XX
XX 05-SEP-2002 (first entry)
XX
XX Murine Toll-like receptor related CpG DNA SEQ ID No 68.
XX
XX Murine Toll-like receptor; TLR9; TLR7; TLR8; ISNA; ds.
XX Unidentified.
XX
XX WO200222809-A2.
XX
XX 21-MAR-2002.
XX
XX 17-SEP-2001; 2001WO-US29229.
XX
XX 15-SEP-2000; 2000US-233035P.
XX 23-JAN-2001; 2001US-263657P.
XX 17-MAY-2001; 2001US-291726P.
XX 22-JUN-2001; 2001US-300210P.
XX
XX (COLE-) COLEY PHARM GMBH.
XX
XX Bauer S, Lipford G, Wagner H;
XX WPI; 2002-393964/42.
XX
XX New isolated murine Toll-like receptor (TLR)9, TLR7, TLR8 polypeptides,
XX useful for identifying species specificity of immunostimulatory nucleic
XX acid and identifying immunostimulatory nucleic acids
XX
XX Disclosure; Page 76; 195pp; English.
XX
XX The invention relates to isolated murine Toll-like receptors (TLR)9,
XX TLR7 and TLR8 polypeptides. These polypeptides comprise fully defined

CC sequences of 1032, 1050 or 1032 amino acids as given in specification, or
CC their fragments, where TLR9, TLR7 and TLR8 polypeptides or their
CC fragments have an amino acid sequence which is identical to human TLR9,
CC TLR7 or TLR8 polypeptides or their fragment except for at least one amino
CC acid of a murine TLR polypeptide. The isolated nucleic acids of the
CC invention are useful for inhibiting TLR9 signalling activity in a cell.
CC TLR7, TLR8 and TLR9 polypeptides are useful for identifying nucleic acid
CC molecules which interact with a TLR polypeptide or its fragment. The
CC TLR7, TLR8 and TLR9 polypeptides are also useful for identifying ISNA. The
CC signalling activity of a test compound (that is not a nucleic acid, and
CC is a polypeptide or a part of a combinatorial library of compounds) with
CC an ISNA. The TLR7, TLR8 and TLR9 polypeptides are also useful for
CC identifying species specificity of an ISNA. The isolated nucleic acids of
CC the invention are useful as probes or primers. This polynucleotide
CC sequence represents DNA relating to the isolated toll-like receptors of
CC the invention.
XX
XX Sequence 20 BP; 5 A; 3 C; 4 G; 8 T; 0 other;
XX
XX Query Match 100.0%; Score 20; DB 24; Length 20;
XX Best Local Similarity 100.0%; Pred. No. 1.3;
XX Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY 1 TATGCATATTCCTGTAAGTG 20
Db | | | | | | | | | | | | | | | | | | | |
1 TATGCATATTCCTGTAAGTG 20
XX
XX
XX RESULT 3
XX AEN80818
XX ID AEN80818 standard; DNA; 24 BP.
XX AC AEN80818;
XX
XX 15-JUL-2002 (first entry)
XX
XX Human STAT-1 inhibitor oligonucleotide SEQ ID NO 36.
XX
XX Human; IRF-1; transcription factor; interferon regulatory factor;
XX antisense; gene therapy; cardiovascular; transplant rejection;
XX immunological hypersensitivity; asthma; inflammatory disease; psoriasis;
XX Crohn's disease; autoimmune disease; diabetes mellitus;
XX multiple sclerosis; rheumatoid arthritis; Th1 response; Th2 response;
XX vasotropic; immunosuppressive; antiasthmatic; dermatological;
XX anti-allergic; anti-ulcer; anti-inflammatory; antipsoriatic; antidiabetic;
XX neuroprotective; antineumatic; antiarthritic; ss.
XX
XX Homo sapiens.
XX Synthetic.
XX
XX WO200229044-A2.
XX
XX 11-APR-2002.
XX
XX 04-OCT-2001; 2001WO-DE03835.
XX
XX 06-OCT-2000; 2000DE-1049549.
XX 29-NOV-2000; 2000DE-1059144.
XX
XX (HECK/) HECKER M.
XX (WAGN/) WAGNER A H.
XX
XX Hecker M, Wagner AH;
XX WPI; 2002-383335/41.
XX
XX Inhibitor of the transcription factor IFR-1, useful for treating e.g.
XX transplant rejection and autoimmune disease, reduces expression of CD40
XX
XX Example 3; Page 19; 45pp; German.
XX

CC The invention relates to an inhibitor (I) of the expression and/or
 CC activity of the transcription factor (IRF-1; interferon regulatory
 CC factor) as a therapeutic agent, especially an oligonucleotide inhibitor
 CC (ABN80783-ABN80804) or antisense oligonucleotide (ABN80805-ABN80808) used
 CC in antisense gene therapy. (I) are used to prevent or treat
 CC cardiovascular complications such as restenosis after angioplasty or
 CC stenosis of venous by-passes, chronic or acute transplant rejection and
 CC graft versus host disease, immunological hypersensitivity, e.g. bronchial
 CC asthma or atopic dermatitis, inflammatory diseases such as ulcerative
 CC colitis, psoriasis and Crohn's disease and autoimmune diseases such as
 CC diabetes mellitus, multiple sclerosis, collagenosis (e.g. systemic lupus
 CC erythematosus), rheumatoid arthritis and vasculitis. (I) simultaneously
 CC weaken both Th1 and Th2 responses. The present sequence is that of an
 CC oligonucleotide, useful to the invention.

XX
 SQ Sequence 24 BP; 6 A; 4 C; 4 G; 10 T; 0 other;

Query Match 95.0%; Score 19; DB 24; Length 24;
 Best Local Similarity 100.0%; Pred. No. 4.3;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAGT 19
 |||||
 Db 6 TATGCATATTCCTGTAAGT 24

RESULT 4
 AAT41604
 ID AAT41604 standard; DNA; 19 BP.

XX
 AC AAT41604;

XX
 DT 04-JUN-1997 (first entry)

XX
 DE Oligonucleotide containing core DNA regulatory element.

XX
 KW Regulatory element; STAT; protein; cytokine; responsive;
 KW host cell; transfection; agonist; antagonist; mediated; STAT5;
 KW transcription; modulation; signalling pathway; STAT6;
 KW oligonucleotide; electrophoretic mobility shift assay; EMSA; ds.

XX
 OS Synthetic.

XX
 FH Key Location/Qualifiers
 FT misc_feature 5..19
 FT /*tag= a
 FT /note= "core DNA regulatory element"

XX
 PN WO9630515-A1.

XX
 PD 03-OCT-1996.

XX
 PF 25-MAR-1996; 96WO-0504012.

XX
 PR 27-MAR-1995; 95US-0411020.

XX
 PA (LIGA-) LIGAND PHARM INC.

XX
 PI Lamb IP, Seidel HM, Tian Chan S;

XX
 DR WPI; 1996-455362/45.

XX
 PT DNA construct for screening modulators of cytokine-mediated
 PT transcription - contg. regulatory element and a cytokine-sensitive
 PT promoter operably linked to a heterologous gene

XX
 PS Example 1; Page 25; 72pp; English.

XX
 CC A novel DNA construct comprises the present oligonucleotide (ON),
 CC which contains a core a regulatory element, operably linked to a
 CC promoter, which is operably linked to a heterologous gene
 CC (preferably a marker gene). The gene is under the transcriptional
 CC control of the promoter and the ON sequence when the ON is bound by

CC a STAT protein activated in response to IL-2, IL-3, G-CSF, GM-CSF,
 CC erythropoietin, thrombopoietin, or preferably IL-4, IL-7, IL-9,
 CC IL-13 or IL-15. Cytokine responsive host cells transfected with the
 CC DNA construct can be used to measure the ability of a compound to
 CC act as an agonist or antagonist of cytokine mediated gene for
 CC transcription. In particular, they can be used to screen for
 CC cytokine modulators involved in the STAT5 and/or STAT6 protein
 CC signalling pathway.
 CC Following an electrophoretic mobility shift assay the DNA construct
 CC was found to bind IL-4 weakly and IL-13 not determined.

XX
 SQ Sequence 19 BP; 5 A; 3 C; 4 G; 7 T; 0 other;

Query Match 80.0%; Score 16; DB 17; Length 19;
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 CATATTCCTGTAAGTG 20
 |||||
 Db 4 CATATTCCTGTAAGTG 19

RESULT 5
 AAT31283
 ID AAT31283 standard; DNA; 17 BP.

XX
 AC AAT31283;

XX
 DT 24-OCT-1996 (first entry)

XX
 DE STAT probe Ly6E.

XX
 KW STAT; signal transducer and activator of transcription;
 KW DNA binding protein; ligand; receptor; oncogenesis; inflammation;
 KW autoimmune disease; antagonist; gene therapy; probe; ds.

XX
 OS Synthetic.

XX
 PN WO9620954-A2.

XX
 PD 11-JUL-1996.

XX
 PF 28-DEC-1995; 95WO-US17025.

XX
 PR 06-JAN-1995; 95US-0369796.

XX
 PA (UYRQ) UNIV ROCKEFELLER.

XX
 PI Darnell JE, Horvath CM, Wen Z, Zhong Z;

XX
 DR WPI; 1996-333941/33.

XX
 PT New STAT protein DNA-binding domain peptide(s) - useful for
 PT diagnosing, preventing or treating cellular dysfunction, e.g.
 PT oncogenesis, inflammation, parasitic disease or autoimmunity

XX
 PS Example 1; Page 42; 138pp; English.

XX
 CC Synthetic DNA probes (AAT31281-86) were used in electrophoretic
 CC mobility shift assays to determine the functionally active
 CC regions of signal transducer and activator of transcription
 CC (STAT) proteins STAT1 (see also AAW03176) and STAT3 (AAW03174).
 CC Amino acids between approx. 400 and approx. 500 of these
 CC human proteins determined DNA binding site specificity.
 CC Mutations within this region resulted in greatly reduced DNA
 CC binding affinities.

XX
 SQ Sequence 17 BP; 5 A; 2 C; 3 G; 7 T; 0 other;

Query Match 75.0%; Score 15; DB 17; Length 17;
 Best Local Similarity 100.0%; Pred. No. 4.2e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATTCTGTAACTG 20
Db 1 ATATTCTGTAACTG 15

RESULT 6

AAQ69287/c
ID AAQ69287 standard; DNA; 47 BP.

XX AC AAQ69287;

XX DT 21-FEB-1995 (first entry)

XX DE Human alpha-amylase gene, target region.

XX KW DNA protein-binding assay; test sequence; screening sequence;
KW promoter; target; TATA box; Herpes Simplex Virus; HSV;
KW origin of replication; UL9; transcription factor; TFIID: ds.

XX OS Synthetic.

XX PN WO9414980-A.

XX PD 07-JUL-1994.

XX PF 20-DEC-1993; 93WO-US12388.

XX PR 23-DEC-1992; 92US-0996783.

XX PR 17-SEP-1993; 93US-0123936.

XX (GENE-) GENELABS TECHNOLOGIES INC.

XX PI Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;

XX WPI; 1994-234711/28.

XX Sequence-directed DNA-binding molecules - useful in
XX pharmaceuticals and as molecular reagents

XX Claim 28; Page 230; 587pp; English.

CC A DNA protein-binding assay is provided, useful for screening
CC libraries of synthetic or biological cpds. for their ability
CC to bind DNA test sequences. The assay is versatile in that any
CC number of test sequences can be tested by placing the test sequence
CC adjacent to a defined protein-binding screening sequence. Binding
CC of mols. to these test sequences changes the binding characteristics
CC of the protein mol. to its cognate binding sequence. When such a mol.
CC binds the test sequence, the equilibrium of the DNA:protein complexes
CC is disturbed, generating changes in the concentration of free DNA probe.
CC One application of this method is to eucaryotic general transcription
CC factors (e.g. TFIID), where the target region is typically selected
CC from DNA sequences adjacent to the binding site for the eucaryotic
CC transcription factor. Numerous exemplary test sequences are given:
CC the sequences in AAQ69251-731 and AAQ69850 correspond to promoter
CC targets (typically, TATA box-contg. sites) for human genes and the
CC sequences in AAQ69732-849 correspond to promoter targets for viral
CC genes. The test sequences may also be randomly generated. DNA:protein
CC interaction may be used for screening purposes, e.g. the Herpes Simplex
CC virus (HSV) origin of replication and UL9 (see AAQ69851-52, AAQ69865 and
CC AAQ69891).

XX SQ Sequence 47 BP; 17 A; 6 C; 10 G; 14 T; 0 other;

Query Match 74.0%; Score 14.8; DB 15; Length 47;
Best Local Similarity 88.9%; Pred. No. 5.9e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18
Db 28 TATTTATATTCCTGTAAG 11

RESULT 7

AAAT63749/c
ID AAT63749 standard; DNA; 47 BP.

XX AC AAT63749;

XX DT 13-MAR-1997 (first entry)

XX DE Human alpha amylase gene TFIID binding site.

XX KW Duplex DNA; target region; binding characteristic; DNA binding protein;
KW TFIID; transcription factor; binding site; inhibition; enhance;
KW cancer; inherited genetic disorder; ds.

XX OS Homo sapiens.

XX PN US5578444-A.

XX PD 26-NOV-1996.

XX PF 27-JUN-1991; 91US-0723618.

XX PR 20-DEC-1993; 93US-0171389.

XX PR 27-JUN-1991; 91US-0723618.

XX PR 23-DEC-1992; 92US-0996783.

XX PR 17-SEP-1993; 93US-0123936.

XX (GENE-) GENELABS TECHNOLOGIES INC.

XX PI Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;

XX WPI; 1997-020402/02.

XX Altering binding characteristics of DNA binding proteins to duplex
XX DNA - by attaching specific small cpd. to target region close to the
XX protein's binding site, useful in treatment of viral disease, cancer
XX etc

XX Claim 6; Column 117; 264pp; English.

CC The sequences given in AAT63713-4312 represent duplex DNA's which act
CC as target regions in the method of the invention. The method for
CC altering the binding characteristics of a DNA-binding protein to duplex
CC DNA comprises contacting the duplex DNA with a small molecule which
CC binds sequence-specifically to a target region, where, when the small
CC molecule is bound to the target region, it is adjacent to, but not
CC overlapping by more than 4 bp, a binding site for a DNA-binding protein.
CC The small molecule is added at a concentration effective to alter the
CC binding of the DNA binding protein, pref. TFIID, to its binding site on
CC the duplex DNA. The binding of the small molecule may inhibit or
CC enhance the binding of the DNA-binding protein to its binding site. The
CC compounds isolated using this method are potentially useful as
CC therapeutic agents for treatment of any disease which involves a
CC specific DNA sequence, e.g. cancer, or inherited genetic disorders etc.
CC The method is suitable for screening large biological or chemical
CC libraries and allows determination of sequence-specific and relative
CC affinities of known DNA-binding agents for different DNA sequences.
CC The design of these duplex DNA's allows a single DNA:protein interaction
CC to be used for screening sequence-specific, or preferential, DNA binding
CC proteins that recognise almost any possible sequence (see also AAT63739-
CC 74).

XX SQ Sequence 47 BP; 17 A; 6 C; 10 G; 14 T; 0 other;

Query Match 74.0%; Score 14.8; DB 18; Length 47;
Best Local Similarity 88.9%; Pred. No. 5.9e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18
Db 28 TATTTATATTCCTGTAAG 11


```

RESULT 8
AAAX17037/c
ID AAX17037 standard; DNA; 47 BP.
XX
AC
XX
AC AAX17037;
XX
DT 06-MAY-1999 (first entry)
XX
XX
DE Test sequence from human alpha-amylase gene.
XX
KW Test sequence; DNA-binding molecule; screening sequence; human;
KW nucleic acid amplification; target; viral; ds.
XX
OS Homo sapiens.
XX
PN US5869241-A.
XX
PD 09-FEB-1999.
XX
PF 07-JUN-1995; 95US-0475228.
XX
PR 20-DEC-1993; 93US-0171389.
PR 27-JUN-1991; 91US-0723618.
PR 23-DEC-1992; 92US-0996783.
PR 17-SEP-1993; 93US-0123936.
PR 07-JUN-1995; 95US-0475228.
XX
PA (GENE-) GENELABS TECHNOLOGIES INC.
XX
PI Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;
XX
WPI; 1999-152755/13.
XX
XX
PT Determination of DNA sequence preference of a DNA-binding molecule -
PT based on inhibition of binding of protein to oligonucleotide
PT sequence attached to test sequence
XX
PS Claim 3; Columns 119-120; 270pp; English.
XX
CC Sequences AAX17001 to AAX17600 represent specifically claimed target
CC test sequences that are used in the method of the invention of
CC determining the DNA sequence preference of a DNA-binding molecule. The
CC method comprises: (i) adding a test molecule and a DNA-binding protein to
CC a mixture of duplex DNA test oligonucleotides, each of the test
CC oligonucleotides having a test sequence adjacent to a screening sequence,
CC where the screening sequence binds to the DNA-binding protein with a
CC binding affinity that is independent of the DNA sequence of the test
CC sequence, and where the mixture of duplex DNA test oligonucleotides
CC includes several test sequences; (ii) incubating the test molecule, the
CC mixture of duplex DNA test oligonucleotides and the DNA-binding protein
CC for a time sufficient to permit binding of the test molecule to test
CC sequences in the duplex DNA; (iii) separating unbound test
CC oligonucleotides from test oligonucleotides bound to binding protein;
CC (iv) amplifying the unbound test oligonucleotides; (v) repeating steps
CC (ii) to (iv); (vi) isolating the amplified test oligonucleotides; and
CC (vii) sequencing the isolated test oligonucleotides. Test sequences
CC AAX17001-X17481 and AAX17600 correspond to promoter targets for human
CC genes and test sequences AAX17482-X17599 correspond to promoter targets
CC for viral genes.
XX
SQ Sequence 47 BP; 17 A; 6 C; 10 G; 14 T; 0 other;
Query Match 74.0%; Score 14.8; DB 20; Length 47;
Best Local Similarity 88.9%; Pred. No. 5.9e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 TATGCATATTCCTGTAAG 18
Db 28 TATTATATTCCTGTAAG 11
RESULT 9
ABK82528/c
ID ABK82528 standard; DNA; 47 BP.
XX
AC
XX
AC ABK82528;
XX
DT 27-AUG-2002 (first entry)
XX
XX
DE DNA binding molecule screening method test sequence #37.
XX
KW DNA binding molecule screening; inhibition of transcription;
KW infection; human immunodeficiency virus; HIV; parasite; cancer;
KW cardiovascular; respiratory; gastrointestinal; endocrine; metabolic;
KW rheumatic; immunological; haematological; neurological;
KW psychiatric; dermatological; ophthalmological; musculo-skeletal;
KW urogenital disorder; ss.
XX
OS Synthetic.
XX
PN US6384208-B1.
XX
PD 07-MAY-2002.
XX
PF 15-JUL-1999; 99US-0354947.
XX
PR 20-DEC-1993; 93US-0171389.
PR 07-JUN-1995; 95US-0482080.
PR 27-JUN-1991; 91US-0723618.
PR 23-DEC-1992; 92US-0996783.
PR 17-SEP-1993; 93US-0123936.
XX
PA (GENE-) GENELABS TECHNOLOGIES INC.
XX
PI Edwards CA, Cantor CR, Andrews BM, Turin LM, Fry KE;
XX
WPI; 2002-442819/47.
XX
XX
PT Decreasing transcriptional activity of genes for treating infections or
PT cancer, by administration of an agent that binds to two non-overlapping
PT regions of the gene
XX
PS Example 15; SEQ ID No 37; 98pp; English.
XX
CC The invention relates to a method of decreasing transcriptional activity
CC in a duplex deoxyribonucleic acid (DNA) template (T1) comprising
CC contacting (T1) with a binding agent comprising at least one small duplex
CC DNA-binding molecule (T2) coupled to at least one other small duplex-
CC binding molecule that binds to a non-overlapping region of target
CC sequence (T3). The method is useful for inhibiting transcription of a
CC range of disease-related genes for treating infections (by viruses,
CC including human immunodeficiency virus, bacteria, fungi, protozoa
CC and parasites), cancer, cardiovascular, respiratory, gastrointestinal,
CC endocrine/metabolic, rheumatic/immunological, haematological,
CC neurological, psychiatric, dermatological, ophthalmological,
CC musculo-skeletal, genetic or urogenital disorders. The method provides
CC sequence-specific inhibition of transcription of pathological genes
CC without affecting transcription of cellular genes regulated by the same
CC transcription factor, and can be applied to regulation of any gene.
CC ABK82492-ABK83155 represent DNA binding molecule test sequences used in
CC the method of the invention.
XX
SQ Sequence 47 BP; 17 A; 6 C; 10 G; 14 T; 0 other;
Query Match 74.0%; Score 14.8; DB 24; Length 47;
Best Local Similarity 88.9%; Pred. No. 5.9e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 TATGCATATTCCTGTAAG 18
Db 28 TATTATATTCCTGTAAG 11
RESULT 10
AAQ69686/c
ID AAQ69686 standard; DNA; 50 BP.

```

XX AAQ69686;
 AC
 XX
 DT
 XX
 XX 02-MAR-1995 (first entry)
 DE Human pancreatic alpha-amylase gene, target region.
 XX
 XX
 KW DNA protein-binding assay; test sequence; screening sequence;
 KW promoter; target; TATA box; Herpes Simplex Virus; HSV;
 KW origin of replication; UL9; transcription factor; TFIID: ds.
 XX
 XX Synthetic.
 OS
 XX
 PN WO9414980-A.
 XX
 PD 07-JUL-1994.
 XX
 XX 20-DEC-1993; 93WO-US12388.
 PF
 XX 23-DEC-1992; 92US-0996783.
 PR
 XX 17-SEP-1993; 93US-0123936.
 XX
 PA (GENE-) GENELABS TECHNOLOGIES INC.
 XX
 XX Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;
 PI
 XX WPI; 1994-234711/28.
 DR
 XX
 XX Sequence-directed DNA-binding molecules - useful in
 PT pharmaceuticals and as molecular reagents
 XX
 XX Claim 28; Page 430; 587pp; English.
 PS
 XX A DNA protein-binding assay is provided, useful for screening
 CC libraries of synthetic or biological cpds. for their ability
 CC to bind DNA test sequences. The assay is versatile in that any
 CC number of test sequences can be tested by placing the test sequence
 CC adjacent to a defined protein-binding screening sequence. Binding
 CC of mols. to these test sequences changes the binding characteristics
 CC of the protein mol. to its cognate binding sequence. When such a mol.
 CC binds the test sequence, the equilibrium of the DNA:protein complexes
 CC is disturbed, generating changes in the concentration of free DNA probe.
 CC One application of this method is to eucaryotic general transcription
 CC factors (e.g. TFIID), where the target region is typically selected
 CC from DNA sequences adjacent to the binding site for the eucaryotic
 CC transcription factor. Numerous exemplary test sequences are given:
 CC the sequences in AAQ69251-731 and AAQ69850 correspond to promoter
 CC targets (typically, TATA box-contg. sites) for human genes and the
 CC sequences in AAQ69732-849 correspond to promoter targets for viral
 CC genes. The test sequences may also be randomly generated. DNA:protein
 CC interaction may be used for screening purposes, e.g. the Herpes Simplex
 CC Virus (HSV) origin of replication and UL9 (see AAQ69851-52, AAQ69865 and
 CC AAQ69891).
 XX
 SQ Sequence 50 BP; 18 A; 6 C; 10 G; 16 T; 0 other;
 Query Match 74.0%; Score 14.8; DB 15; Length 50;
 Best Local Similarity 88.9%; Pred. No. 5.9e+02;
 Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 TATGCATATTCCTGTAAG 18
 ||| |||||
 DB 32 TATTATATTCCTGTAAG 15
 RESULT 11
 AAQ69687/c
 ID AAQ69687 standard; DNA; 50 BP.
 XX
 AC AAQ69687;
 XX
 DT 02-MAR-1995 (first entry)
 XX

DE
 XX Human pancreatic alpha-amylase gene, target region.
 KW DNA protein-binding assay; test sequence; screening sequence;
 KW promoter; target; TATA box; Herpes Simplex Virus; HSV;
 KW origin of replication; UL9; transcription factor; TFIID: ds.
 XX
 XX Synthetic.
 OS
 XX
 PN WO9414980-A.
 XX
 PD 07-JUL-1994.
 XX
 XX 20-DEC-1993; 93WO-US12388.
 PF
 XX 23-DEC-1992; 92US-0996783.
 PR
 XX 17-SEP-1993; 93US-0123936.
 XX
 PA (GENE-) GENELABS TECHNOLOGIES INC.
 XX
 XX Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;
 PI
 XX WPI; 1994-234711/28.
 DR
 XX
 XX Sequence-directed DNA-binding molecules - useful in
 PT pharmaceuticals and as molecular reagents
 XX
 XX Claim 28; Page 430; 587pp; English.
 PS
 XX A DNA protein-binding assay is provided, useful for screening
 CC libraries of synthetic or biological cpds. for their ability
 CC to bind DNA test sequences. The assay is versatile in that any
 CC number of test sequences can be tested by placing the test sequence
 CC adjacent to a defined protein-binding screening sequence. Binding
 CC of mols. to these test sequences changes the binding characteristics
 CC of the protein mol. to its cognate binding sequence. When such a mol.
 CC binds the test sequence, the equilibrium of the DNA:protein complexes
 CC is disturbed, generating changes in the concentration of free DNA probe.
 CC One application of this method is to eucaryotic general transcription
 CC factors (e.g. TFIID), where the target region is typically selected
 CC from DNA sequences adjacent to the binding site for the eucaryotic
 CC transcription factor. Numerous exemplary test sequences are given:
 CC the sequences in AAQ69251-731 and AAQ69850 correspond to promoter
 CC targets (typically, TATA box-contg. sites) for human genes and the
 CC sequences in AAQ69732-849 correspond to promoter targets for viral
 CC genes. The test sequences may also be randomly generated. DNA:protein
 CC interaction may be used for screening purposes, e.g. the Herpes Simplex
 CC Virus (HSV) origin of replication and UL9 (see AAQ69851-52, AAQ69865 and
 CC AAQ69891).
 XX
 SQ Sequence 50 BP; 18 A; 6 C; 10 G; 16 T; 0 other;
 Query Match 74.0%; Score 14.8; DB 15; Length 50;
 Best Local Similarity 88.9%; Pred. No. 5.9e+02;
 Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 TATGCATATTCCTGTAAG 18
 ||| |||||
 DB 32 TATTATATTCCTGTAAG 15
 RESULT 12
 AAT64148/c
 ID AAT64148 standard; DNA; 50 BP.
 XX
 AC AAT64148;
 XX
 DT 17-MAR-1997 (first entry)
 XX
 DE Human pancreatic alpha-amylase gene TFIID binding site.
 XX
 KW Duplex DNA; target region; binding characteristic; DNA binding protein;
 KW TFIID; transcription factor; binding site; inhibition; enhance;
 KW cancer; inherited genetic disorder; ds.

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XX OS Homo sapiens.
XX OS
XX PN US5578444-A.
XX PN
XX PD 26-NOV-1996.
XX PD
XX PF 27-JUN-1991; 91US-0723618.
XX PF
XX PR 20-DEC-1993; 93US-0171389.
XX PR
XX PR 27-JUN-1991; 91US-0723618.
XX PR
XX PR 23-DEC-1992; 92US-0996783.
XX PR
XX PR 17-SEP-1993; 93US-0123936.
XX PR
XX PA (GENE-) GENELABS TECHNOLOGIES INC.
XX PA
XX PI Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;
XX PI
XX DR WPI; 1997-020402/02.
XX DR
XX PT Altering binding characteristics of DNA binding proteins to duplex
XX PT DNA - by attaching specific small cpd. to target region close to the
XX PT protein's binding site, useful in treatment of viral disease, cancer
XX PT etc
XX PS Claim 6; Column 321-322; 264pp; English.
XX PS
XX CC The sequences given in AAT63713-4312 represent duplex DNA's which act
XX CC as target regions in the method of the invention. The method for
XX CC altering the binding characteristics of a DNA-binding protein to duplex
XX CC DNA comprises contacting the duplex DNA with a small molecule which
XX CC binds sequence-specifically to a target region, where, when the small
XX CC molecule is bound to the target region, it is adjacent to, but not
XX CC overlapping by more than 4 bp, a binding site for a DNA-binding protein.
XX CC The small molecule is added at a concentration effective to alter the
XX CC binding of the DNA binding protein, pref. TFIID, to its binding site on
XX CC the duplex DNA. The binding of the small molecule may inhibit or
XX CC enhance the binding of the DNA-binding protein to its binding site. The
XX CC compounds isolated using this method are potentially useful as
XX CC therapeutic agents for treatment of any disease which involves a
XX CC specific DNA sequence, e.g. cancer, or inherited genetic disorders etc.
XX CC The method is suitable for screening large biological or chemical
XX CC libraries and allows determination of sequence-specific and relative
XX CC affinities of known DNA-binding agents for different DNA sequences.
XX CC The design of these duplex DNA's allows a single DNA:protein interaction
XX CC to be used for screening sequence-specific, or preferential, DNA binding
XX CC proteins that recognise almost any possible sequence (see also AAT49539-
XX CC 74).
XX CC
XX SQ Sequence 50 BP; 18 A; 6 C; 10 G; 16 T; 0 other;
XX SQ
XX Query Match 74.0%; Score 14.8; DB 18; Length 50;
XX Best Local Similarity 88.9%; Pred. No. 5.9e+02;
XX Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
XX
XX QY 1 TATGCATATTCCTGTAAG 18
XX ||| |||||
XX Db 32 TATTTATATTCCTGTAAG 15
XX
XX RESULT 13
XX AAT64149/c
XX ID AAT64149 standard; DNA; 50 BP.
XX AC
XX AC AAT64149;
XX XX
XX DT 17-MAR-1997 (first entry)
XX DT
XX DE Human pancreatic amylase gene TFIID binding site.
XX DE
XX KW Duplex DNA; target region; binding characteristic; DNA binding protein;
XX KW TFIID; transcription factor; binding site; inhibition; enhance;
XX KW cancer; inherited genetic disorder; ds.
XX KW

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XX OS Homo sapiens.
XX OS
XX PN US5578444-A.
XX PN
XX PD 26-NOV-1996.
XX PD
XX PF 27-JUN-1991; 91US-0723618.
XX PF
XX PR 20-DEC-1993; 93US-0171389.
XX PR
XX PR 27-JUN-1991; 91US-0723618.
XX PR
XX PR 23-DEC-1992; 92US-0996783.
XX PR
XX PR 17-SEP-1993; 93US-0123936.
XX PR
XX PA (GENE-) GENELABS TECHNOLOGIES INC.
XX PA
XX PI Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;
XX PI
XX DR WPI; 1997-020402/02.
XX DR
XX PT Altering binding characteristics of DNA binding proteins to duplex
XX PT DNA - by attaching specific small cpd. to target region close to the
XX PT protein's binding site, useful in treatment of viral disease, cancer
XX PT etc
XX PS Claim 6; Column 321-322; 264pp; English.
XX PS
XX CC The sequences given in AAT63713-4312 represent duplex DNA's which act
XX CC as target regions in the method of the invention. The method for
XX CC altering the binding characteristics of a DNA-binding protein to duplex
XX CC DNA comprises contacting the duplex DNA with a small molecule which
XX CC binds sequence-specifically to a target region, where, when the small
XX CC molecule is bound to the target region, it is adjacent to, but not
XX CC overlapping by more than 4 bp, a binding site for a DNA-binding protein.
XX CC The small molecule is added at a concentration effective to alter the
XX CC binding of the DNA binding protein, pref. TFIID, to its binding site on
XX CC the duplex DNA. The binding of the small molecule may inhibit or
XX CC enhance the binding of the DNA-binding protein to its binding site. The
XX CC compounds isolated using this method are potentially useful as
XX CC therapeutic agents for treatment of any disease which involves a
XX CC specific DNA sequence, e.g. cancer, or inherited genetic disorders etc.
XX CC The method is suitable for screening large biological or chemical
XX CC libraries and allows determination of sequence-specific and relative
XX CC affinities of known DNA-binding agents for different DNA sequences.
XX CC The design of these duplex DNA's allows a single DNA:protein interaction
XX CC to be used for screening sequence-specific, or preferential, DNA binding
XX CC proteins that recognise almost any possible sequence (see also AAT49539-
XX CC 74).
XX CC
XX SQ Sequence 50 BP; 18 A; 6 C; 10 G; 16 T; 0 other;
XX SQ
XX Query Match 74.0%; Score 14.8; DB 18; Length 50;
XX Best Local Similarity 88.9%; Pred. No. 5.9e+02;
XX Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
XX
XX QY 1 TATGCATATTCCTGTAAG 18
XX ||| |||||
XX Db 32 TATTTATATTCCTGTAAG 15
XX
XX RESULT 14
XX AAX17436/c
XX ID AAX17436 standard; DNA; 50 BP.
XX AC
XX AC AAX17436;
XX XX
XX DT 06-MAY-1999 (first entry)
XX DT
XX DE Test sequence from human pancreatic alpha-amylase gene.
XX DE
XX KW Test sequence; DNA-binding molecule; screening sequence; human;
XX KW nucleic acid amplification; target; viral; ds.
XX KW

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OS Homo sapiens.
XX US5869241-A.
XX
XX
XX
XX
XX
XX PF 07-JUN-1995; 950S-0475228.
XX
XX 20-DEC-1993; 930S-0171389.
XX 27-JUN-1991; 910S-0723618.
XX 23-DEC-1992; 920S-0996783.
XX 17-SEP-1993; 930S-0123936.
XX 07-JUN-1995; 950S-0475228.
XX
XX (GENE-) GENELABS TECHNOLOGIES INC.
XX
XX Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;
XX
XX WPI; 1999-152755/13.
XX
XX Determination of DNA sequence preference of a DNA-binding molecule -
XX based on inhibition of binding of protein to oligonucleotide
XX sequence attached to test sequence
XX
XX Claim 3; Columns 323-324; 270pp: English.
XX
XX Sequences AAX17001 to AAX17600 represent specifically claimed target
XX test sequences that are used in the method of the invention of
XX determining the DNA sequence preference of a DNA-binding molecule. The
XX method comprises: (i) adding a test molecule and a DNA-binding protein to
XX a mixture of duplex DNA test oligonucleotides, each of the test
XX oligonucleotides having a test sequence adjacent to a screening sequence,
XX where the screening sequence binds to the DNA-binding protein with a
XX binding affinity that is independent of the DNA sequence of the test
XX sequence, and where the mixture of duplex DNA test oligonucleotides
XX includes several test sequences; (ii) incubating the test molecule, the
XX mixture of duplex DNA test oligonucleotides and the DNA-binding protein
XX for a time sufficient to permit binding of the test molecule to test
XX sequences in the duplex DNA; (iii) separating unbound test
XX oligonucleotides from test oligonucleotides bound to binding protein;
XX (iv) amplifying the unbound test oligonucleotides; (v) repeating steps
XX (ii) to (iv); (vi) isolating the amplified test oligonucleotides; and
XX (vii) sequencing the isolated test oligonucleotides. Test sequences
XX AAX17001-X17481 and AAX17600 correspond to promoter targets for human
XX genes and test sequences AAX17482-X17599 correspond to promoter targets
XX for viral genes.
XX
XX Sequence 50 BP; 18 A; 6 C; 10 G; 16 T; 0 other;
XX
XX Query Match 74.0%; Score 14.8; DB 20; Length 50;
XX Best Local Similarity 88.9%; Pred. No. 5.9e+02;
XX Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
XX
XX QY 1 TATGCATATTCCTGTAAG 18
XX ||| |||||
XX
XX Db 32 TATTTATATTCCTGTAAG 15
XX
XX
XX RESULT 15
XX AAX17437/c
XX ID AAX17437 standard; DNA; 50 BP.
XX
XX AC AAX17437;
XX
XX DT 06-MAY-1999 (first entry)
XX
XX DE Test sequence from human pancreatic amylase gene.
XX
XX KW Test sequence; DNA-binding molecule; screening sequence; human;
XX nucleic acid amplification; target; viral; ds.
XX
XX OS Homo sapiens.
XX

```

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PN US5869241-A.
XX
XX
XX
XX
XX
XX PF 07-JUN-1995; 950S-0475228.
XX
XX 20-DEC-1993; 930S-0171389.
XX 27-JUN-1991; 910S-0723618.
XX 23-DEC-1992; 920S-0996783.
XX 17-SEP-1993; 930S-0123936.
XX 07-JUN-1995; 950S-0475228.
XX
XX (GENE-) GENELABS TECHNOLOGIES INC.
XX
XX Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;
XX
XX WPI; 1999-152755/13.
XX
XX Determination of DNA sequence preference of a DNA-binding molecule -
XX based on inhibition of binding of protein to oligonucleotide
XX sequence attached to test sequence
XX
XX Claim 3; Columns 323-324; 270pp: English.
XX
XX Sequences AAX17001 to AAX17600 represent specifically claimed target
XX test sequences that are used in the method of the invention of
XX determining the DNA sequence preference of a DNA-binding molecule. The
XX method comprises: (i) adding a test molecule and a DNA-binding protein to
XX a mixture of duplex DNA test oligonucleotides, each of the test
XX oligonucleotides having a test sequence adjacent to a screening sequence,
XX where the screening sequence binds to the DNA-binding protein with a
XX binding affinity that is independent of the DNA sequence of the test
XX sequence, and where the mixture of duplex DNA test oligonucleotides
XX includes several test sequences; (ii) incubating the test molecule, the
XX mixture of duplex DNA test oligonucleotides and the DNA-binding protein
XX for a time sufficient to permit binding of the test molecule to test
XX sequences in the duplex DNA; (iii) separating unbound test
XX oligonucleotides from test oligonucleotides bound to binding protein;
XX (iv) amplifying the unbound test oligonucleotides; (v) repeating steps
XX (ii) to (iv); (vi) isolating the amplified test oligonucleotides; and
XX (vii) sequencing the isolated test oligonucleotides. Test sequences
XX AAX17001-X17481 and AAX17600 correspond to promoter targets for human
XX genes and test sequences AAX17482-X17599 correspond to promoter targets
XX for viral genes.
XX
XX Sequence 50 BP; 18 A; 6 C; 10 G; 16 T; 0 other;
XX
XX Query Match 74.0%; Score 14.8; DB 20; Length 50;
XX Best Local Similarity 88.9%; Pred. No. 5.9e+02;
XX Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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XX QY 1 TATGCATATTCCTGTAAG 18
XX ||| |||||
XX
XX Db 32 TATTTATATTCCTGTAAG 15
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XX Job time : 105.319 secs

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GenCore version 5.1.3
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:06:16 ; Search time 318.116 Seconds
(without alignments)
1829.698 Million cell updates/sec

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Perfect score: 20
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Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 2054640 segs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0
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Post-processing: Minimum Match 0%
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Listing first 45 summaries

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- 2: gb_hgt.*
- 3: gb_in.*
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- 37: em_htg_vrt.*
- 38: em_sy.*
- 39: em_htgo_hum.*
- 40: em_htgo_mus.*
- 41: em_htgo_other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	20	100.0	20	6	A89787 Sequence 9
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3	20	100.0	20	6	AR078393 Sequence 9
4	20	100.0	20	6	AX455575 Sequence
5	20	100.0	27	6	AR063217 Sequence
6	20	100.0	27	6	AX001643 Sequence
7	20	100.0	27	6	I49528 Sequence 2
8	19	95.0	20	6	A89784 Sequence 6
9	19	95.0	20	6	A90871 Sequence 6
10	16.4	82.0	32	6	AR135319 Sequence
11	16.4	82.0	50	6	AR032839 Sequence
12	16.4	82.0	50	6	AR209503 Sequence
13	16.4	82.0	50	6	I29579 Sequence 45
14	16.4	82.0	50	6	I91253 Sequence 45
15	15.8	79.0	20	6	A89786 Sequence 8
16	15.8	79.0	20	6	A90873 Sequence 8
17	14.4	72.0	83	6	AX328437 Sequence
18	14.2	71.0	41	6	AR109119 Sequence
19	14.2	71.0	41	6	AR200774 Sequence
20	13.8	69.0	71	6	AR122803 Sequence
21	13.6	68.0	51	6	AX162422 Sequence
22	13.2	66.0	25	6	AX117480 Sequence
23	13.2	66.0	51	6	AX117481 Sequence
24	13.2	66.0	64	9	AF032251 Otolenur
25	13.2	66.0	75	9	D86841 Homo sapien
26	13.2	66.0	76	9	AF032248 Otolenur
27	13.2	66.0	76	9	AF032254 Otolenur
28	13	65.0	26	6	A89779 Sequence 1
29	13	65.0	26	6	A90893 Sequence 28
30	12.8	64.0	21	6	A91594 Sequence 12
31	12.8	64.0	51	6	AX159923 Sequence
32	12.8	64.0	51	6	AX159925 Sequence
33	12.8	64.0	51	6	AX159926 Sequence
34	12.8	64.0	51	6	AX160807 Sequence
35	12.8	64.0	51	6	AX160808 Sequence
36	12.8	64.0	51	6	AX164818 Sequence
37	12.8	64.0	51	6	AX189849 Sequence
38	12.8	64.0	61	6	AR118244 Sequence
39	12.8	64.0	76	9	AF032239 Otolenur
40	12.8	64.0	81	11	G34985 BPY1 human
41	12.6	63.0	57	6	AR064089 Sequence
42	12.6	63.0	57	6	AR064090 Sequence
43	12.6	63.0	57	6	BD008488 Targeting
44	12.6	63.0	57	6	BD008489 Targeting
45	12.6	63.0	58	6	AX053230 Sequence

ALIGNMENTS

RESULT 1	A89787	Sequence 9 from Patent WO9832462.	20 bp	DNA	linear	PAT 22-JAN-2000
LOCUS	A89787	Sequence 9 from Patent WO9832462.				
DEFINITION	A89787					
ACCESSION	A89787.1	GI:6738301				
VERSION						
KEYWORDS						
SOURCE						
ORGANISM						
REFERENCE	1	(bases 1 to 20)				
AUTHORS	Lipford,G.B. and Heeg,K.					
TITLE	PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND					
JOURNAL	OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION					
	Patent: WO 9832462-A 9 30-JUL-1998;					

● **●**

VERSION AX001643.1 GI:7241772
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 27)
AUTHORS Damert,A. and Plate,K.
TITLE REGULATORY SEQUENCES INVOLVED IN HYPOXIA REGULATED GENE EXPRESSION
AND USES THEREOF
JOURNAL Patent: WO 9856936-A 15 17-DEC-1998;
MAX PLANCK GESSELLSCHAFT (DE); DAMERT ANNETTE (DE)
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Db 4 GATTGCTGACGTCAGAG 23

RESULT 7
LOCUS 149528 27 bp DNA linear PAT 07-OCT-1997
DEFINITION Sequence 2 from patent US 5641486.
ACCESSION 149528
VERSION 149528.1 GI:2471748
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 27)
AUTHORS Hintichs,S.H. and Orten,D.Jo.
TITLE Methods for inhibiting transcription of the cyclic AMP responsive
element binding protein and the activating transcription factor 1
JOURNAL Patent: US 5641486-A 2 24-JUN-1997;
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source Location/Qualifiers
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/organism="unknown"
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAG 20
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Db 4 GATTGCTGACGTCAGAG 23

RESULT 8
LOCUS A89784 20 bp DNA linear PAT 22-JAN-2000
DEFINITION Sequence 6 from Patent WO9832462.
ACCESSION A89784
VERSION A89784.1 GI:6738298
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Lipford,G.B. and Heeg,K.
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
JOURNAL Patent: WO 9832462-A 6 30-JUL-1998;
LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)

FEATURES
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Db 1 ATTGCCTGACGTCAGAG 19

RESULT 9
LOCUS A90871 20 bp DNA linear PAT 22-JAN-2000
DEFINITION Sequence 6 from Patent EP0855184.
ACCESSION A90871
VERSION A90871.1 GI:6739265
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Heeg,K.P. and Lipford,G.B.
TITLE Pharmaceutical composition comprising a polynucleotide and an
antigen especially for vaccination
JOURNAL Patent: EP 0855184-A 6 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
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Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 ATTGCCTGACGTCAGAG 20
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Db 1 ATTGCCTGACGTCAGAG 19

RESULT 10
LOCUS ARI35319 32 bp DNA linear PAT 16-MAY-2001
DEFINITION Sequence 1 from patent US 6194632.
ACCESSION ARI35319
VERSION ARI35319.1 GI:14124224
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 32)
AUTHORS Leiden,J.M.
TITLE Mouse model for congestive heart failure
JOURNAL Patent: US 6194632-A 1 27-FEB-2001;
FEATURES
source Location/Qualifiers
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QY 3 TTGCCTGACGTCAGAG 20

Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ATTGCCTGACGTCAGAG 20
 Db 1 ATTGCCTGACGTCAGAG 19

Search completed: December 12, 2002, 02:55:19
 Job time : 326.116 secs

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GenCore version 5.1.3
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OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:06:16 ; Search time 318.116 Seconds
(without alignments)
1829.698 Million cell updates/sec

Title: US-09-355-254F-21

Perfect score: 20
Sequence: 1 gtatttcacagaaaggaac 20

Scoring table: IDENTITY_NUC
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Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600.

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Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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18: em_in.*
19: em_mu.*
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36: em_hcg_mam.*
37: em_hcg_vrt.*
38: em_sy.*
39: em_hgo_hum.*
40: em_hgo_mus.*
41: em_hgo_other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	20	100.0	20	6	A89800	A89800 Sequence 22
2	20	100.0	20	6	A90887	A90887 Sequence 22
3	20	100.0	20	6	AR010171	AR010171 Sequence
4	20	100.0	20	6	AR204918	AR204918 Sequence
5	20	100.0	20	6	AR204923	AR204923 Sequence
6	20	100.0	20	6	AX455586	AX455586 Sequence
7	20	100.0	20	6	I33466	I33466 Sequence 3
8	20	100.0	20	6	I40053	I40053 Sequence 3
9	20	100.0	20	6	I47062	I47062 Sequence 3
10	20	100.0	20	6	I81487	I81487 Sequence 3
11	20	100.0	30	6	AR182577	AR182577 Sequence 3
12	20	100.0	42	6	A37863	A37863 Sequence 6
13	20	100.0	100	6	AR174601	AR174601 Sequence
14	20	100.0	100	6	AR174602	AR174602 Sequence
15	20	100.0	100	6	AX047032	AX047032 Sequence
16	20	100.0	100	6	AX047033	AX047033 Sequence
17	20	100.0	100	6	AX280202	AX280202 Sequence
18	20	100.0	100	6	AX280203	AX280203 Sequence
19	20	100.0	100	6	AX365191	AX365191 Sequence
20	20	100.0	100	6	AX365192	AX365192 Sequence
21	17	85.0	17	6	AR165233	AR165233 Sequence
22	17	85.0	17	6	AR201404	AR201404 Sequence
23	17	85.0	17	6	AX459960	AX459960 Sequence
24	17	85.0	17	6	I87794	I87794 Sequence 22
25	16	80.0	25	6	AR182576	AR182576 Sequence
26	16	80.0	25	6	AR083256	AR083256 Sequence
27	16	80.0	25	6	AR182574	AR182574 Sequence
28	16	80.0	30	6	AR048146	AR048146 Sequence
29	15.8	79.0	90	9	HSEXTR2	X99574 H.sapiens e
30	15	75.0	15	6	AX338644	AX338644 Sequence
31	15	75.0	15	6	I15312	I15312 Sequence 13
32	15	75.0	15	6	I31724	I31724 Sequence 13
33	15	75.0	15	6	I93568	I93568 Sequence 13
34	14.2	71.0	29	11	DOGCFTRC	L77689 Canis famli
35	14.2	71.0	39	10	MMVCAM1C7	U12877 Mus musculu
36	14.2	71.0	57	9	HSAPOLPA1	AF139730 Homo sapi
37	14.2	71.0	69	3	AF250305	AF250305 Chironomu
38	14.2	71.0	98	6	AX354846	AX354846 Sequence
39	14	70.0	16	6	AR165232	AR165232 Sequence
40	14	70.0	16	6	AR201403	AR201403 Sequence
41	14	70.0	16	6	AX338643	AX338643 Sequence
42	14	70.0	16	6	AX459959	AX459959 Sequence
43	14	70.0	16	6	I15311	I15311 Sequence 12
44	14	70.0	16	6	I31723	I31723 Sequence 12
45	14	70.0	16	6	I93567	I93567 Sequence 12

ALIGNMENTS

RESULT 1	A89800	Sequence 22 from Patent WO9832462.	20 bp	DNA	linear	PAT 22-JAN-2000
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DEFINITION	A89800	Sequence 22 from Patent WO9832462.				
ACCESSION	A89800	Sequence 22 from Patent WO9832462.				
VERSION	A89800.1	GI:6738314				
KEYWORDS		unidentified.				
SOURCE		unidentified.				
ORGANISM		unclassified.				
REFERENCE		1 (bases 1 to 20)				
AUTHORS		Lipford,G.B. and Heeg,K.				
TITLE		PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND				
JOURNAL		OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION				
		Patent: WO 9832462-A 22 30-JUL-1998;				

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Db	1	GTATTTCCCGAGAAAGGAAC	20
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LOCUS	AR204918	20 bp	DNA linear PAT 20-JUN-2002
DEFINITION	Sequence 9 from patent US 6368828.		
ACCESSION	AR204918		
VERSION	AR204918.1	GI:21502366	
KEYWORDS	unknown.		
SOURCE	Unknown.		
ORGANISM	Unclassified.		
REFERENCE	1 (bases 1 to 20)		
AUTHORS	Laroche,W., Patel,B.K.R. and Pierce,J.H.		
TITLE	Attenuated and dominant negative variant cDNAs of Stat6; Stat6b and Stat6c		
JOURNAL	Patent: US 6368828-A 9 09-APR-2002;		
FEATURES	Location/Qualifiers		
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Db	1	GTATTTCCCGAGAAAGGAAC	20
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DEFINITION	Sequence 14 from patent US 6368828.		
ACCESSION	AR204923		
VERSION	AR204923.1	GI:21502372	
KEYWORDS	unknown.		
SOURCE	Unknown.		
ORGANISM	Unclassified.		
REFERENCE	1 (bases 1 to 20)		
AUTHORS	Laroche,W., Patel,B.K.R. and Pierce,J.H.		
TITLE	Attenuated and dominant negative variant cDNAs of Stat6; Stat6b and Stat6c		
JOURNAL	Patent: US 6368828-A 14 09-APR-2002;		
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Matches	20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;		
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Db	1	GTATTTCCCGAGAAAGGAAC	20
RESULT 6			
LOCUS	AX455586	20 bp	DNA linear PAT 06-JUL-2002
DEFINITION	Sequence 63 from Patent WO0222809.		
ACCESSION	AX455586		
VERSION	AX455586.1	GI:21714654	
KEYWORDS			

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SOURCE      synthetic construct.
ORGANISM     synthetic construct
artificial sequences.
REFERENCE    1
AUTHORS      Bauer,S., Lipford,G. and Wagner,H.
TITLE        Process for high throughput screening of cpq-based
              immuno-agonist/antagonist
JOURNAL      Patent: WO 0222809-A 63 21-MAR-2002;
              Coley Pharmaceutical GmbH (DE)
FEATURES     Location/Qualifiers
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RESULT 7
I33466
LOCUS          I33466          20 bp      DNA      linear      PAT 06-FEB-1997
DEFINITION     Sequence 3 from patent US 5591825.
ACCESSION      I33466
VERSION        I33466.1 GI:1824257
KEYWORDS       Unknown.
SOURCE         Unknown.
ORGANISM       Unclassified.
REFERENCE      1 (bases 1 to 20)
AUTHORS        McKnight,S.L. and Hou,J.
TITLE          Interleukin 4 signal transducers
JOURNAL        Patent: US 5591825-A 3 07-JAN-1997;
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    |||||
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LOCUS          I40053          20 bp      DNA      linear      PAT 13-MAY-1997
DEFINITION     Sequence 3 from patent US 5618693.
ACCESSION      I40053
VERSION        I40053.1 GI:2083058
KEYWORDS       Unknown.
SOURCE         Unknown.
ORGANISM       Unclassified.
REFERENCE      1 (bases 1 to 20)
AUTHORS        McKnight,S.L., Hou,J. and Schindler,U.
TITLE          Interleukin-2 signal transducers and binding assays
JOURNAL        Patent: US 5618693-A 3 08-APR-1997;
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DEFINITION     Sequence 3 from patent US 5639858.
ACCESSION      I47062
VERSION        I47062.1 GI:2471027
KEYWORDS       Unknown.
SOURCE         Unknown.
ORGANISM       Unclassified.
REFERENCE      1 (bases 1 to 20)
AUTHORS        Hoeft,T. and Rothe,M.
TITLE          Human signal transducer and binding assays
JOURNAL        Patent: US 5639858-A 3 17-JUN-1997;
FEATURES       Location/Qualifiers
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Db 1 GTATTTCCCGAGAAAGGAAC 20

RESULT 10
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LOCUS          I81487          20 bp      DNA      linear      PAT 10-JUN-1998
DEFINITION     Sequence 3 from patent US 5710266.
ACCESSION      I81487
VERSION        I81487.1 GI:3209784
KEYWORDS       Unknown.
SOURCE         Unknown.
ORGANISM       Unclassified.
REFERENCE      1 (bases 1 to 20)
AUTHORS        McKnight,S.L. and Hou,J.
TITLE          Nucleic acid encoding an interleukin 4 signal transducer
JOURNAL        Patent: US 5710266-A 3 20-JAN-1998;
FEATURES       Location/Qualifiers
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RESULT 11
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LOCUS          AR182577          30 bp      DNA      linear      PAT 20-APR-2002
DEFINITION     Sequence 25 from patent US 6338949.

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ACCESSION AR182577
VERSION AR182577.1 GI:20225784
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 30)
AUTHORS Darnell, J.E. Jr., Schindler, C.W., Fu, X.-Y., Wen, Z. and Zhong, Z.
TITLE Nucleic acids encoding receptor recognition factor stat4 and methods of use thereof
JOURNAL Patent: US 6338949-A 25 15-JAN-2002;
FEATURES Location/Qualifiers
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BASE COUNT 10 a 8 c 6 g 6 t
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RESULT 12
A37863
LOCUS A37863
DEFINITION Sequence 6 from Patent WO9408025.
ACCESSION A37863
VERSION A37863.1 GI:2294543
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 42)
AUTHORS Benach, P., Perez, C. and Wietzerbin, J.
TITLE DNA SEQUENCES INVOLVED IN THE TRANSCRIPTION OF GENES UNDER THE EFFECT OF INDUCERS, AND BIOLOGICAL APPLICATIONS THEREOF
JOURNAL Patent: WO 9408025-A 6 14-APR-1994;
COMMENT INST NAT SANTE RECH MED (FR)
OTHER PUBLICATION FR 2696181 940401.
FEATURES Location/Qualifiers
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/db_xref="taxon:32644"
BASE COUNT 14 a 6 c 10 g 12 t
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 23 GTATTTCAGAAAGGAAC 42

RESULT 13
AR174601
LOCUS AR174601
DEFINITION Sequence 59 from patent US 6307024.
ACCESSION AR174601
VERSION AR174601.1 GI:17914921
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 100)
AUTHORS Novak, J.E., Presnell, S.R., Sprecher, C.A., Foster, D.C., Holly, R.D., Gross, J.A., Johnston, J.V., Nelson, A.J., Dillon, S.R. and Hammond, A.K.

QY 1 GTATTTCAGAAAGGAAC 20
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Db 44 GTATTTCAGAAAGGAAC 63

RESULT 14
AR174602/c
LOCUS AR174602
DEFINITION Sequence 60 from patent US 6307024.
ACCESSION AR174602
VERSION AR174602.1 GI:17914922
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 100)
AUTHORS Novak, J.E., Presnell, S.R., Sprecher, C.A., Foster, D.C., Holly, R.D., Gross, J.A., Johnston, J.V., Nelson, A.J., Dillon, S.R. and Hammond, A.K.
TITLE Cytokine zalphall Ligand
JOURNAL Patent: US 6307024-A 60 23-OCT-2001;
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BASE COUNT 26 a 17 c 32 g 25 t
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LOCUS AX047032
DEFINITION Sequence 37 from Patent WO0068381.
ACCESSION AX047032
VERSION AX047032.1 GI:11876456
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 100)
AUTHORS Presnell, S.R., Foster, D.C., Hammond, A.K. and Lok, S.
TITLE Cytokine receptor mouse zcytor10
JOURNAL Patent: WO 0068381-A 37 16-NOV-2000;
ZymoGenetics, Inc. (US)
FEATURES Location/Qualifiers
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/db_xref="taxon:32630"
/note="Oligonucleotide primer ZC12749"
BASE COUNT 24 a 33 c 17 g 26 t
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTATTTCACAGAAAGGAC 20
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Db 44 GTATTTCACAGAAAGGAC 63

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Job time : 322.116 secs

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Run on: December 12, 2002, 01:06:16 ; Search time 318.116 Seconds
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1829.698 Million cell updates/sec

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Perfect score: 20
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Gapop 10.0 , Gapext 1.0

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Post-processing: Minimum Match 0%
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- 40: em.htgo.mus.*
- 41: em.htgo.other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
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2	20	100.0	20	6	A90886	A90886 Sequence 21
3	20	100.0	20	6	AX455553	AX455553 Sequence 21
4	20	100.0	21	6	I23456	I23456 Sequence 8
5	20	100.0	100	6	ARI74601	ARI74601 Sequence 8
c 6	20	100.0	100	6	ARI74602	ARI74602 Sequence
c 7	20	100.0	100	6	AX047032	AX047032 Sequence
c 8	20	100.0	100	6	AX047033	AX047033 Sequence
c 9	20	100.0	100	6	AX280202	AX280202 Sequence
c 10	20	100.0	100	6	AX280203	AX280203 Sequence
c 11	20	100.0	100	6	AX365191	AX365191 Sequence
c 12	20	100.0	100	6	AX365192	AX365192 Sequence
c 13	14.2	71.0	31	6	ARI19638	ARI19638 Sequence
c 14	14.2	71.0	31	6	ARI169119	ARI169119 Sequence
c 15	14.2	71.0	58	6	ARI38229	ARI38229 Sequence
c 16	14.2	71.0	64	12	SYNCOCK	K01192 Yeast (S.ce
c 17	13.8	69.0	24	6	A03717	A03717 Oligonucleo
c 18	13.8	69.0	41	6	AX343814	AX343814 Sequence
c 19	13.8	69.0	41	6	AX343816	AX343816 Sequence
c 20	13.8	69.0	44	6	AX343812	AX343812 Sequence
c 21	13.8	69.0	44	6	AX343818	AX343818 Sequence
c 22	13.8	69.0	76	8	YSCGNC152	M87429 Yeast Eco R
c 23	13.8	69.0	91	1	AF087321	AF087321 Chlamydia
c 24	13.8	69.0	94	10	AY041972	AY041972 Phodopus
c 25	13.6	68.0	25	6	AR043941	AR043941 Sequence
c 26	13.6	68.0	25	6	AR073474	AR073474 Sequence
c 27	13.6	68.0	25	6	AX022068	AX022068 Sequence
c 28	13.6	68.0	25	6	I93345	I93345 Sequence 15
c 29	13.6	68.0	26	6	AR043992	AR043992 Sequence
c 30	13.6	68.0	26	6	AR073525	AR073525 Sequence
c 31	13.6	68.0	26	6	AX022135	AX022135 Sequence
c 32	13.6	68.0	26	6	I93396	I93396 Sequence 82
c 33	13.6	68.0	65	6	AX483015	AX483015 Sequence
c 34	13.6	68.0	72	6	AR043980	AR043980 Sequence
c 35	13.6	68.0	72	6	AR073513	AR073513 Sequence
c 36	13.6	68.0	72	6	AX022123	AX022123 Sequence
c 37	13.6	68.0	72	6	I93384	I93384 Sequence 70
c 38	13.2	56.0	31	6	AX327657	AX327657 Sequence
c 39	13.2	56.0	45	6	AR076832	AR076832 Sequence
c 40	13.2	56.0	51	6	AX160107	AX160107 Sequence
c 41	13.2	56.0	58	6	ARI40194	ARI40194 Sequence
c 42	13.2	56.0	58	6	ARI73275	ARI73275 Sequence
c 43	13.2	56.0	63	6	ARI38221	ARI38221 Sequence
c 44	13.2	56.0	72	6	I74780	I74780 Sequence 12
c 45	13.2	56.0	74	10	RATPAM25	U52661 Rattus norv

ALIGNMENTS

RESULT 1
A89799 A89799 Sequence 21 from Patent WO9832462. 20 bp DNA linear PAT 22-JAN-2000
LOCUS
DEFINITION
ACCESSION A89799
VERSION
KEYWORDS
SOURCE unidentified.
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Lipford G.B. and Heeg K.
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
JOURNAL Patent: WO 9832462-A 21 30-JUL-1998;

FEATURES
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LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
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BASE COUNT 7 a 3 c 3 g 7 t
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Best Local Similarity 100.0%; Pred. No. 5.5e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGATTCTAGGAATTCATC 20
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Db 1 AGATTCTAGGAATTCATC 20
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LOCUS A90886 20 bp DNA linear PAT 22-JAN-2000
DEFINITION Sequence 21 from Patent EP0855184.
ACCESSION A90886
VERSION A90886.1 GI:6739328
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified
REFERENCE 1 (bases 1 to 20)
AUTHORS Heeg,K.P. and Lipford,G.B.
TITLE Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination
JOURNAL Patent: EP 0855184-A 21 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
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BASE COUNT 7 a 3 c 3 g 7 t
ORIGIN

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Best Local Similarity 100.0%; Pred. No. 5.5e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGATTCTAGGAATTCATC 20
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Db 1 AGATTCTAGGAATTCATC 20
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RESULT 3
LOCUS AX45553 20 bp DNA linear PAT 06-JUL-2002
DEFINITION Sequence 30 from Patent WO0222809.
ACCESSION AX45553
VERSION AX45553.1 GI:21714621
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1
AUTHORS Bauer,S., Lipford,G. and Wagner,H.
TITLE process for high throughput screening of cpg-based immuno-agonist/antagonist
JOURNAL Patent: WO 0222809-A 30 21-MAR-2002;
Coley Pharmaceutical GmbH (DE)
FEATURES
source
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/note="synthetic oligonucleotide"
BASE COUNT 7 a 3 c 3 g 7 t
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RESULT 4
LOCUS I23456 21 bp DNA linear PAT 07-OCT-1996
DEFINITION Sequence 8 from patent US 5534409.
ACCESSION I23456
VERSION I23456.1 GI:1603326
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 21)
AUTHORS Groner,B., Gouilleux,F. and Wakao,H.
TITLE Cytokine regulated transcription factor
JOURNAL Patent: US 5534409-A 8 09-JUL-1996;
FEATURES
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BASE COUNT 7 a 4 c 3 g 7 t
ORIGIN

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Best Local Similarity 100.0%; Pred. No. 5.4e+02;
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Db 1 AGATTCTAGGAATTCATC 20
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RESULT 5
LOCUS ARI74601 100 bp DNA linear PAT 17-DEC-2001
DEFINITION Sequence 59 from patent US 6307024.
ACCESSION ARI74601
VERSION ARI74601.1 GI:17914921
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 100)
AUTHORS Novak,J.E., Presnell,S.R., Sprecher,C.A., Foster,D.C., Holly,R.D., Gross,J.A., Johnston,J.V., Nelson,A.J., Dillon,S.R. and Hammond,A.K.
TITLE Cytokine zaiphal1 Ligand
JOURNAL Patent: US 6307024-A 59 23-OCT-2001;
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BASE COUNT 25 a 32 c 17 g 26 t
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Db 67 AGATTCTAGGAATTCATC 86
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RESULT 6
LOCUS ARI74602/c 100 bp DNA linear PAT 17-DEC-2001
DEFINITION Sequence 60 from patent US 6307024.

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ACCESSION   AR174602
VERSION     AR174602.1  GI:17914922
KEYWORDS    Unknown.
SOURCE      Unknown.
ORGANISM    Unclassified.
REFERENCE   1 (bases 1 to 100)
AUTHORS    Novak,J.E., Presnell,S.R., Sprecher,C.A., Foster,D.C., Holly,R.D.,
            Gross,J.A., Johnston,J.V., Nelson,A.J., Dillon,S.R. and
            Hammond,A.K.
TITLE      Cytokine zalphall Ligand
JOURNAL    Patent: US 6307024-A 60 23-OCT-2001;
FEATURES    Location/Qualifiers
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RESULT 7
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LOCUS       AX047032             100 bp      DNA      linear      PAT 15-DEC-2000
DEFINITION Sequence 37 from Patent WO0068381.
ACCESSION   AX047032
VERSION     AX047032.1  GI:11876456
KEYWORDS    synthetic construct.
SOURCE      synthetic construct.
ORGANISM    artificial sequences.
REFERENCE   1 (bases 1 to 100)
AUTHORS    Presnell,S.R., Foster,D.C., Hammond,A.K. and Lok,S.
TITLE      Cytokine receptor mouse zcytor10
JOURNAL    Patent: WO 0068381-A 37 16-NOV-2000;
            ZymoGenetics, Inc. (US)
FEATURES    Location/Qualifiers
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BASE COUNT  24 a 33 c 17 g 26 t
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Db   67 AGATTTCTAGGAATTC AATC 86

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LOCUS       AX047033             100 bp      DNA      linear      PAT 15-DEC-2000
DEFINITION Sequence 38 from Patent WO0068381.
ACCESSION   AX047033
VERSION     AX047033.1  GI:11876457
KEYWORDS    synthetic construct.
SOURCE      synthetic construct.
ORGANISM    artificial sequences.
REFERENCE   1 (bases 1 to 100)
AUTHORS    Presnell,S.R., Foster,D.C., Hammond,A.K. and Lok,S.
TITLE      Cytokine receptor mouse zcytor10
JOURNAL    Patent: WO 0068381-A 38 16-NOV-2000;

ACCESSION   AR174602
VERSION     AR174602.1  GI:17914922
KEYWORDS    Unknown.
SOURCE      Unknown.
ORGANISM    Unclassified.
REFERENCE   1 (bases 1 to 100)
AUTHORS    Novak,J.E., Presnell,S.R., Sprecher,C.A., Foster,D.C., Holly,R.D.,
            Gross,J.A., Johnston,J.V., Nelson,A.J., Dillon,S.R. and
            Hammond,A.K.
TITLE      Cytokine zalphall Ligand
JOURNAL    Patent: US 6307024-A 60 23-OCT-2001;
FEATURES    Location/Qualifiers
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BASE COUNT  26 a 17 c 32 g 25 t
ORIGIN

Query Match      100.0%; Score 20; DB 6; Length 100;
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Db   38 AGATTTCTAGGAATTC AATC 19

RESULT 9
AX280202
LOCUS       AX280202             100 bp      DNA      linear      PAT 02-NOV-2001
DEFINITION Sequence 48 from Patent WO0177171.
ACCESSION   AX280202
VERSION     AX280202.1  GI:16607595
KEYWORDS    synthetic construct.
SOURCE      synthetic construct.
ORGANISM    artificial sequences.
REFERENCE   1
AUTHORS    Sprecher,C.A., Novak,J.E., West,J.W., Presnell,S.R., Holly,R.D. and
            Nelson,A.J.
TITLE      Soluble zalphall cytokine receptors
JOURNAL    Patent: WO 0177171-A 48 18-OCT-2001;
            ZymoGenetics, Inc. (US)
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Db   67 AGATTTCTAGGAATTC AATC 86

RESULT 10
AX280203/c
LOCUS       AX280203             100 bp      DNA      linear      PAT 02-NOV-2001
DEFINITION Sequence 49 from Patent WO0177171.
ACCESSION   AX280203
VERSION     AX280203.1  GI:16607596
KEYWORDS    synthetic construct.
SOURCE      synthetic construct.
ORGANISM    artificial sequences.
REFERENCE   1
AUTHORS    Sprecher,C.A., Novak,J.E., West,J.W., Presnell,S.R., Holly,R.D. and
            Nelson,A.J.
TITLE      Soluble zalphall cytokine receptors
JOURNAL    Patent: WO 0177171-A 49 18-OCT-2001;
            ZymoGenetics, Inc. (US)
FEATURES    Location/Qualifiers
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               /db_xref="taxon:32630"
               /note="Oligonucleotide primer ZC12748"
BASE COUNT  26 a 17 c 33 g 24 t
ORIGIN
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AUTHORS Sheppard, P.O. and Humes, J.M.
 TITLE Adipocyte-specific protein homologs
 JOURNAL Patent: US 6197930-A 46 06-MAR-2001;
 FEATURES Location/Qualifiers
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 BASE COUNT 17 a 16 c 9 g 16 t
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 Db 14 AGAATACTAGGAATTCAT 32

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 Job time : 322.116 secs

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GenCore version 5.1.3
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OM nucleic - nucleic search, using sw model

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Perfect score: 20
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Gapop 10.0, Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600

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Maximum Match 100%
Listing first 45 summaries

Database :

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17: em_hum:*
18: em_in:*
19: em_mu:*
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27: em_sts:*
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29: em_vl:*
30: em_htg_hum:*
31: em_htg_inv:*
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score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
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2	20	100.0	20	6	A90885	A90885 Sequence 20
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c 4	20	100.0	21	6	I39725	I39725 Sequence 12
c 5	20	100.0	21	6	I55842	I55842 Sequence 12
6	18	90.0	86	6	AR184059	AR184059 Sequence
7	18	90.0	86	6	AR203344	AR203344 Sequence
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10	15.2	76.0	24	6	AR061864	AR061864 Sequence
11	15	75.0	19	6	AR043736	AR043736 Sequence
c 12	15	75.0	19	6	AR043737	AR043737 Sequence
13	15	75.0	19	6	I81940	I81940 Sequence 38
c 14	15	75.0	19	6	I81941	I81941 Sequence 39
c 15	13.8	69.0	27	3	AF251713S4	AF251713S4
c 16	13.8	69.0	27	3	AF251718S4	AF251718S4
c 17	13.8	69.0	27	3	AF251743S4	AF251743S4
18	13.8	69.0	65	6	AX485415	AX485415 Sequence
c 19	13.6	68.0	41	6	AX088050	AX088050 Sequence
c 20	13.6	68.0	51	6	AX115257	AX115257 Sequence
c 21	13.6	68.0	100	1	ECOFIMPHC	M11776 E.coli inve
22	13.4	67.0	20	6	AR208137	AR208137 Sequence
c 23	13.4	67.0	60	6	AR009405	AR009405 Sequence
c 24	13.4	67.0	90	3	PFASSRR	D17580 Plasmidium
c 25	13.4	67.0	90	6	AR022104	AR022104 Sequence
c 26	13.4	67.0	90	6	AR022120	AR022120 Sequence
c 27	13.4	67.0	90	6	E08156	E08156 Primer or p
28	13.2	66.0	28	6	I39716	I39716 Sequence 3
29	13.2	66.0	28	6	I55833	I55833 Sequence 3
30	13.2	66.0	35	6	A92293	A92293 Sequence 12
31	13.2	66.0	35	6	A92344	A92344 Sequence 12
32	13.2	66.0	35	6	BD007234	BD007234 Lentiviru
c 33	13.2	66.0	38	6	I15748	I15748 Sequence 14
c 34	13.2	66.0	41	6	AX088048	AX088048 Sequence
c 35	13.2	66.0	48	6	AX429820	AX429820 Sequence
c 36	13.2	66.0	83	12	AY096764	AY096764 Populus t
37	13.2	66.0	90	10	RAT11BHVSD	M77835 R.norvegicu
38	13	65.0	13	6	AX026535	AX026535 Sequence
39	12.8	64.0	20	6	AX296543	AX296543 Sequence
40	12.8	64.0	24	6	AX291910	AX291910 Sequence
c 41	12.8	64.0	27	3	AF251728S4	AF251731 Dicyrtoma
c 42	12.8	64.0	27	3	AF251738S4	AF251741 Bilobella
c 43	12.8	64.0	48	9	HS4308535	AJ308535 Homo sapi
c 44	12.8	64.0	49	6	A43525	A43525 Sequence 5
c 45	12.8	64.0	49	6	AR052464	AR052464 Sequence

ALIGNMENTS

RESULT 1

A89798
LOCUS A89798
DEFINITION Sequence 20 from Patent WO9832462.
ACCESSION A89798
VERSION A89798.1 GI:6738312
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL

A89798
Sequence 20 from Patent WO9832462.
A89798
A89798.1 GI:6738312
unidentified.
unidentified.
unclassified.
1 (bases 1 to 20)
Lipford,G.B. and Heeg,K.
PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
Patent: WO 9832462-A 20 30-JUL-1998;

20 bp
DNA
linear
PAT 22-JAN-2000

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FEATURES
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LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
Location/Qualifiers
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BASE COUNT 5 a 5 c 4 g 6 t
ORIGIN

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Db 1 CTGATTCCCGGAATGATG 20

RESULT 2
A90885
LOCUS A90885 20 bp DNA linear PAT 22-JAN-2000
DEFINITION Sequence 20 from Patent EP0855184.
ACCESSION A90885
VERSION A90885.1 GI:6739323
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Heeg,K.P. and Lipford,G.B.
TITLE Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination
JOURNAL Patent: EP 0855184-A 20 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
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RESULT 3
AX455567
LOCUS AX455567 20 bp DNA linear PAT 06-JUL-2002
DEFINITION Sequence 44 from Patent WO0222809.
ACCESSION AX455567
VERSION AX455567.1 GI:21714635
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1
AUTHORS Bauer,S., Lipford,G. and Wagner,H.
TITLE Process for high throughput screening of cpq-based immuno-agonist/antagonist
JOURNAL Patent: WO 0222809-A 44 21-MAR-2002;
Coley Pharmaceutical GmbH (DE)
FEATURES
source Location/Qualifiers
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/note="Synthetic oligonucleotide"
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Db 1 CTGATTCCCGGAATGATG 20

RESULT 4
I39725
LOCUS I39725 21 bp DNA linear PAT 13-MAY-1997
DEFINITION Sequence 12 from patent US 5616489.
ACCESSION I39725
VERSION I39725.1 GI:2084205
KEYWORDS
SOURCE unknown.
ORGANISM unknown.
REFERENCE 1 (bases 1 to 21)
AUTHORS Levy,D.E.
TITLE DNA sequence which binds transcriptional regulatory proteins activated in response to various cytokines and uses thereof
JOURNAL Patent: US 5616489-A 12 01-APR-1997;
FEATURES
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1. .21
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Db 20 CTGATTCCCGGAATGATG 1

RESULT 5
I55842
LOCUS I55842 21 bp DNA linear PAT 07-OCT-1997
DEFINITION Sequence 12 from patent US 5648217.
ACCESSION I55842
VERSION I55842.1 GI:2476636
KEYWORDS
SOURCE unknown.
ORGANISM unknown.
REFERENCE 1 (bases 1 to 21)
AUTHORS Levy,D.E.
TITLE DNA sequence which binds transcriptional regulatory proteins activated in response to various cytokines and uses thereof
JOURNAL Patent: US 5648217-A 12 15-JUL-1997;
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RESULT 6
AR184059
LOCUS AR184059 86 bp DNA linear PAT 20-APR-2002
DEFINITION Sequence 3 from patent US 6342581.


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ACCESSION AR184059
VERSION AR184059.1 GI:20228028
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 86)
AUTHORS Rosen,C.A., Ruben,S.M., Olsen,H.S. and Ebner,R.
TITLE Secreted protein HLHP03
JOURNAL Patent: US 6342581-A 3 29-JAN-2002;
FEATURES
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ORIGIN

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Db 40 TGATTTCCTCCCGAAATGAT 57
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DEFINITION Sequence 6 from patent US 6365369.
ACCESSION AR203344
VERSION AR203344.1 GI:21499709
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 86)
AUTHORS Endress,G.A. and Rosen,C.A.
TITLE Prostate specific secreted protein
JOURNAL Patent: US 6365369-A 6 02-APR-2002;
FEATURES
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QY 2 TGATTTCCTCCCGAAATGAT 19
    |||||||
Db 40 TGATTTCCTCCCGAAATGAT 57
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RESULT 8
LOCUS AR206966
DEFINITION Sequence 18 from patent US 6372473.
ACCESSION AR206966
VERSION AR206966.1 GI:21505728
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 86)
AUTHORS Moore,P.A., Ruben,S.M. and Ebner,R.
TITLE Tissue plasminogen activator-like protease
JOURNAL Patent: US 6372473-A 18 16-APR-2002;
FEATURES
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ORIGIN

Query Match
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Db 40 TGATTTCCTCCCGAAATGAT 57
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RESULT 9
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DEFINITION Sequence 24 from patent US 5843697.
ACCESSION AR062163
VERSION AR062163.1 GI:5989854
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 22)
AUTHORS Pestka,S. and Kotenko,S.V.
TITLE Cells expressing IL-10 receptor and the CRFB4 gene product, an IL-10 receptor accessory protein
JOURNAL Patent: US 5843697-A 24 01-DEC-1998;
FEATURES
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RESULT 10
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DEFINITION Sequence 56 from patent US 5843660.
ACCESSION AR061864
VERSION AR061864.1 GI:5989555
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 24)
AUTHORS Schumm,J.W., Micka,K.A. and Rabbach,D.R.
TITLE Multiplex amplification of short tandem repeat loci
JOURNAL Patent: US 5843660-A 56 01-DEC-1998;
FEATURES
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BASE COUNT 5 a 6 c 4 g 9 t
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RESULT 11
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DEFINITION Sequence 106 from patent US 5814517.
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ACCESSION AR043736
 VERSION AR043736.1 GI:5964744
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 UNCLASSIFIED.
 REFERENCE 1 (bases 1 to 19)
 AUTHORS Seidel,H.Martin., and Lamb,I.Peter.
 TITLE DNA spacer regulatory elements responsive to cytokines and methods for their use
 JOURNAL Patent: US 5814517-A 106 29-SEP-1998;
 FEATURES Location/Qualifiers
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 BASE COUNT 5 a 5 c 4 g 5 t
 ORIGIN

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QY 3 GATTTCCTCCCGAAATG 17
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 Db 5 GATTTCCTCCCGAAATG 19
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RESULT 12
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 LOCUS AR043737 19 bp DNA linear PAT 29-SEP-1999
 DEFINITION Sequence 107 from patent US 5814517.
 ACCESSION AR043737
 VERSION AR043737.1 GI:5964745
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 UNCLASSIFIED.
 REFERENCE 1 (bases 1 to 19)
 AUTHORS Seidel,H.Martin., and Lamb,I.Peter.
 TITLE DNA spacer regulatory elements responsive to cytokines and methods for their use
 JOURNAL Patent: US 5814517-A 107 29-SEP-1998;
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 BASE COUNT 5 a 4 c 5 g 5 t
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QY 3 GATTTCCTCCCGAAATG 17
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 Db 5 GATTTCCTCCCGAAATG 19
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RESULT 13
 I81940
 LOCUS I81940 19 bp DNA linear PAT 10-JUN-1998
 DEFINITION Sequence 38 from patent US 5712094.
 ACCESSION I81940
 VERSION I81940.1 GI:3210237
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 UNCLASSIFIED.
 REFERENCE 1 (bases 1 to 19)
 AUTHORS Seidel,H.Martin., Lamb,I.Peter., and Chan,S.-S.Tian.
 TITLE Methods for detecting modulators of cytokine action
 JOURNAL Patent: US 5712094-A 38 27-JAN-1998;
 FEATURES Location/Qualifiers
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BASE COUNT 5 a 5 c 4 g 5 t
 ORIGIN

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 Best Local Similarity 100.0%; Pred. No. 2.1e+03;
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QY 3 GATTTCCTCCCGAAATG 17
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 Db 5 GATTTCCTCCCGAAATG 19
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 LOCUS I81941 19 bp DNA linear PAT 10-JUN-1998
 DEFINITION Sequence 39 from patent US 5712094.
 ACCESSION I81941
 VERSION I81941.1 GI:3210238
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 UNCLASSIFIED.
 REFERENCE 1 (bases 1 to 19)
 AUTHORS Seidel,H.Martin., Lamb,I.Peter., and Chan,S.-S.Tian.
 TITLE Methods for detecting modulators of cytokine action
 JOURNAL Patent: US 5712094-A 39 27-JAN-1998;
 FEATURES Location/Qualifiers
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 Db 19 GATTTCCTCCCGAAATG 5
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RESULT 15
 AF251713S4/c
 LOCUS AF251713S4 27 bp DNA linear INV 12-OCT-2000
 DEFINITION Acерentomon sp. CFNDS1 elongation factor 1-alpha gene, exon 4.
 ACCESSION AF251716
 VERSION AF251716.1 GI:10798923
 KEYWORDS
 SEGMENT 4 of 5
 SOURCE Acерentomon sp. CFNDS1.
 ORGANISM Acерentomon sp. CFNDS1
 Eukaryota; Metazoa; Arthropoda; Hexapoda; Protura; Acерentomidae;
 Acерentomidae; Acерentomon.
 REFERENCE 1 (bases 1 to 27)
 AUTHORS Carapelli,A., Frati,F., Nardi,F., Dallai,R. and Simon,C.
 TITLE Molecular phylogeny of the Apterygotan insects based on nuclear and mitochondrial genes
 JOURNAL Pedobiologia (Jena) 44, 361-373 (2000)
 REFERENCE 2 (bases 1 to 27)
 AUTHORS Carapelli,A., Frati,F., Nardi,F., Dallai,R. and Simon,C.
 TITLE Direct Submission
 JOURNAL Submitted (30-MAR-2000) Evolutionary Biology, University of Siena, via P.A. Mattioli 4, Siena 53100, Italy
 COMMENT Region: Presence of introns of unknown length between s 1&2, 2&3, 3&4.
 Region: The five s joined together correspond to bases 2108 to 2617 in Drosophila melanogaster F1 sequence (accession X06869).
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Db 19 TGATTTCCTCGAAACGA 3

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